



Environment and Time as Constraints on the Biogeographical Distribution of Gibbons

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Highlights:

- Time budgets limit the biogeographic distribution of gibbons and siamang
- Time that has to be allocated to resting seems to be the main constraint
- The limits on group size are not so restrictive as to make monogamy obligatory

For Peer Review

**Environment and Time as Constraints on the Biogeographical
Distribution of Gibbons**

by

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Abstract

We develop a time budget model for the hylobatid family with the aim of assessing the extent to which their contemporary and historical biogeographic distributions might be explained by ecological constraints. The model uses local climate to predict time budgets, and from this the limiting size of social group that animals could manage at a given location. The model predicts maximum group sizes that vary between 3-15 within the taxon’s current distribution, indicating that the combination of their dietary and locomotor styles with the kinds of habitats they inhabit radically constrain group size. Beyond the edges of their current distribution, sustainable group size rapidly tends to zero, although if they had been able to bypass some of these areas, they would have found very suitable habitats in southern India and beyond the Wallace Line. While travel time would be a major constraint on group size at larger group sizes, as it is in great apes, the main factor limiting the gibbon’s current distribution is the time they need to spend resting that is imposed on them by the environment. The model also indicates that gibbons would not now be able to survive in regions of central and southeastern China where they are known to have occurred within historical times, perhaps because historical climate change following the Little Ice Age of the C18th made these regions uninhabitable for them. Finally, our results indicate that gibbons have the ecological capacity to live in larger groups than they do, making it unlikely that their adoption of monogamy reflects purely ecological constraints.

Key Words: climate, biogeographic distribution, group size, foraging ecology

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- Time that has to be allocated to resting seems to be the main constraint
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Introduction

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64 Inability to satisfy nutritional and other demands within a defined time period
65 (usually a 24-hr physiological cycle: Peters 1983) limits both where a species can live
66 and, in social species, how big their groups can be (Dunbar, Korstjens & Lehmann
67 2009). In this respect, time is a central problem for all animals, especially for those
68 that, like primates, are not active 24 hours a day. This insight has led to the
69 development of a series of time budget models designed to understand the role of time
70 in limiting the biogeographical distributions of individual taxa (Dunbar et al. 2009).
71 These models are premised on the fact that behaviour (as the outcome of core
72 physiological processes) is the interface between the environment and the animals’
73 ability to survive (with the latter indexed by the size of group they can maintain as an
74 ecological entity) (see also Marshall, Carter, Rowcliffe & Cowlishaw 2013). There is
75 little that most animals can do to alter their physiology in response to changes in
76 climate or vegetation, but they can, and do, adjust their behaviour and group sizes in
77 response to changing environmental conditions.

78 For species like anthropoid primates that (with obvious the exception *Aotus*)
79 are strictly diurnal due to poor night vision, the time available for satisfying their
80 nutritional requirements is further reduced to the ~12 hours of tropical daylight. In
81 addition, climate and the species’ dietary adaptations may force animals to rest
82 (‘enforced rest’ *sensu* Korstjens et al. 2010), thereby further reducing the length of
83 their active day. The climatic component of this relationship reflects the fact that
84 almost all tropical animals are obliged to cease being active and seek shelter during
85 the hottest part of the day when ambient temperatures in open sunlight exceed their
86 thermoneutral zone (Mount 1979; Dàvid-Barrett & Dunbar 2016); the dietary aspect is
87 mainly a problem for folivores for whom gut fermentation of foliage is incompatible

with any form of activity because the microbial activity on which it depends is extremely sensitive to the heat generated by physical activity (van Soest 1982).

Intensely social species, like most anthropoid primates, face an additional problem because the functionality of their relationships, as well as the cohesion of their social groups, depends directly on the time they invest in grooming with each other (Lehmann, Korstjens & Dunbar 2007a; Pollard & Blumstein 2008; Dunbar & Shultz 2010; Dunbar & Lehmann 2013; Sutcliffe, Dunbar, Binder & Arrow 2012; Dunbar 2018a,b). For both humans and nonhuman primates, an individual's willingness to give coalitionary aid to another depends directly on the amount of time they spend engaged in affiliative interaction (Seyfarth & Cheney 1984; Dunbar 1980, 2018a,b; Burton-Chellew & Dunbar 2015). At some point, animals will inevitably run out of time, and this will ultimately limit where they can live. Ecologists tend to see this as a constraint imposed by nutrient availability, but the real constraint is time.

In effect, animals face a three-way optimization problem in which they have to offset the costs of living in social groups (i.e. time investment in social grooming) and the acquisition of nutrients (specified by the ecological determinants of time required for feeding and travel, including the fixed costs of fuelling fertility and lactation) against the benefits of group size as their principal form of predator deterrence. Predation risk will depend on whether the animal is terrestrial or arboreal and on the hunting styles of the local predators (the 'landscape of fear') (Shultz et al. 2004; Shultz & Finlayson 2010; Tolon et al. 2009; Laundré et al. 2014; Coleman & Hill 2014; Riginos 2015; Gallagher, Creel, Wilson & Cooke 2017), but within these constraints the prey species can adjust group size to offset local predation risk.

Focusing on time budgets is not an alternative to the more conventional socio-ecological approach that typically seeks to identify correlations between, on the one

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hand, group size, population density or behaviour and, on the other hand, climate or forage quality/distribution (Wrangham, Gittleman & Chapman 1993; Chapman & Chapman 2000; Snaith & Chapman 2007; Clutton-Brock & Janson 2012), or approaches that explore the efficiency of foraging (e.g. Brockleman et al. 2014). Rather, a focus on time budgets provides a way of looking at the mechanisms that underpin the correlations between environmental variables and population level outcomes (group size and dispersion). Time budgets thus provide insights into the mechanisms involved in these processes, and, in doing so, emphasize a component of that mechanism (namely, time) that is invariably overlooked in most studies.

In addition, a time budget approach allows us to incorporate, directly or indirectly, a range of other factors and causal relationships that bear on animals' ability to survive in particular locations. This is important because biological phenomena are naturally systems-based, with most causal relationships being multivariate and subject to the influence of feedback loops. Failure to incorporate these more complex relationships may result in the over- or underestimation of the importance of particular causal relationships. Time budget models allow us to integrate a range of variables and relationships into a single, coherent model.

The formal structure of time budget models is that of a causal chain. The climate at a given location directly or (via its effect on vegetation quality) indirectly determines the three core elements of the time budget (feeding, moving and minimum or 'enforced' rest time). These in turn determine the limiting size of group that animals can maintain at that location because this depends on how much 'uncommitted' time the animals then have available and the parametric relationship between time invested in relationships and the resulting quality of those relationships (as reflected in group coherence). The maximum possible group size predicted by the

time budget ultimately determines whether or not the species can live at that location (Dunbar et al. 2009). It is important to appreciate that these models are location-specific: they make explicit predictions about the behaviour of the animals at a given location, subject to the particular climatic and vegetational conditions pertaining at that location. They are also necessarily genus-specific because they reflect the particular dietary, physiological and body size adaptations that characterize a genus. In this respect, a genus is an ecological species.

Climate variables affect both energy demand (energy animals need to maintain thermoregulation: Mount 1979) and the rate of nutrient intake through the effect climate has on the quality and digestibility of forage (and hence the amount of forage that has to be eaten to extract the required amount of nutrients: van Soest 1982). They also influence both travel time (through their effect on foraging patch size and inter-patch distances: Janson & van Schaik 1988; Chapman & Chapman 2000) and rest time (when animals are forced to seek shelter because ambient temperatures exceed their thermoneutral zone: Mount 1979; Dàvid-Barrett & Dunbar 2016). In addition, the feedback loops that are invariably an important component of biological systems can arise when group size, in particular, influences travel time (because larger groups have to travel further if feeding patches are depleted easily – typically, less of a problem for folivores), and this in turn requires more time to be devoted to foraging to replace the energy consumption of the additional travel.

Time budget models consist of a set of simultaneous equations (one for each of the core time budget variables) and thus have the form of a linear programming (or linear optimization) model (Dunbar 2002). The intersection of these equations defines the limit on group size that a species can maintain at a given location. If any of these relationships involve nonlinear components, solving the simultaneous equation set to

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3 163 find the maximum group size usually requires numerical rather than analytical
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5 164 methods (Dunbar 1992a; Dunbar et al. 2009). However, if all relationships are linear,
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7 165 analytical methods based on substitution can be used. The set of climatic parameters
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9 166 (the primary drivers, with each as a separate X, Z,... axis) and group size (the final
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11 167 output, or Y-axis, variable) define a multidimensional state space. The isosurface
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13 168 defined by points where maximum group size drops below the minimum required for
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15 169 demographic viability and/or predator defence defines the taxon's biogeographic
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17 170 distribution. So long as the predicted maximum group size lies above these minima,
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19 171 the species can survive at those locations, with the magnitude of the difference
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21 172 between the maximum and these minima providing a measure of how much
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23 173 ecological stress the animals are under.
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28 174 These mechanism-based models are at least as successful as conventional
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30 175 climate (or niche) envelope models at predicting the geographical distributions of the
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32 176 taxa concerned (Willems & Hill 2009; Korstjens et al. 2010). In addition, they provide
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34 177 insights into which aspects of the animals' ecology are responsible for limiting their
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36 178 distribution (Dunbar et al. 2009). This is because time budget models allow us (a) to
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38 179 determine how much ecological and demographic stress a particular population is
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40 180 under (Dunbar 1992a; Lehmann et al. 2007a) and (b) to specify why the species is
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42 181 unable to occupy particular habitats in ways that directly reflect their physiological
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44 182 adaptations (Dunbar et al. 2009). Neither of these is possible with conventional
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46 183 climate envelope models, which are essentially simple correlational relationships. In
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48 184 addition, because time budget models are driven directly by climate, the models can
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50 185 be used to explore the consequences of past and future climate change for a taxon's
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52 186 biogeographic distribution in a more nuanced way (Lehmann et al. 2010; Kortsjens,
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54 187 Lehmann & Dunbar 2010; Bettridge & Dunbar 2012).
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3 188 Time budget models have been published for a number of terrestrial (gelada:
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5 189 Dunbar 1992a; baboons: Dunbar 1992b; Bettridge, Lehmann & Dunbar 2010; African
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7 190 great apes: Lehmann, Korstjens & Dunbar 2007b, 2008a) and arboreal (spider
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9 191 monkeys: Korstjens, Verhoeckx & Dunbar 2006; colobins: Korstjens & Dunbar 2007;
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11 192 vervets: Willems & Hill 2009; forest guenons: Korstjens, Lehmann & Dunbar 2018;
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13 193 orangutans: Carne, Semple & Lehmann 2012) primates, as well as one ungulate (feral
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15 194 goats: Dunbar & Shi 2013).

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19 195 Although models have been developed for all three great apes (Lehmann et al.
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21 196 2007b, 2008a,b; Carne et al. 2012), the small apes (the gibbon family, Hylobatidae)
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23 197 have yet to be considered. The small apes offer a particularly interesting challenge for
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25 198 several reasons. First, for an ape, they occupy an unusually exclusive arboreal
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27 199 terminal branch niche otherwise occupied only by some of the smaller monkeys.
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29 200 Second, they typically live in very small groups (monogamous pairs plus dependent
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31 201 offspring) and this raises questions as to why they have opted for this form of social
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33 202 system. One classic explanation for monogamy in mammals is that females are forced
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35 203 by their ecology to forage on their own in territories that are too large for a male to be
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37 204 able to successfully defend more than one female (Wittenberg & Tilson 1980; van
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39 205 Schaik & Dunbar 1990; Komers & Brotherton 1997; Lukas & Clutton-Brock 2013);
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41 206 an alternative suggestion is that male parental support allows females to reduce their
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43 207 energy demand, especially in strongly seasonal habitats or where locomotor costs
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45 208 prohibit fat storage (Dunbar 1995a,b; Heldstab, van Schaik & Isler 2017). Third, they
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47 209 have a relatively limited distribution in southeast Asia, and this limited range begs
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56 211 Historically, hylobatids probably evolved as a distinct family in southwest
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213 Malay peninsula into the islands of the Sunda shelf as opportunities provided by
214 changes in climate and sea level allowed (Jablonski & Chaplin 2009). Turvey, Crees
215 & Di Fonzo (2015) found that, as late as the 1800s, gibbons were still being recorded
216 in historical documents as widely distributed throughout southern and central China as
217 far north as latitude ~35°N (Shanxi, Shaanxi and Shangdong provinces). What
218 explains their disappearance from all but the southwest corner of China (Yunnan and
219 Hainan) remains to be determined, though anthropogenic factors have been suspected
220 (Fan 2017).

221 In this paper, we have five main aims. The first is to develop a model of
222 hylobatid time budgets in terms of climatic variables. Second, we test the model's
223 validity by using it to predict the taxon's presence and absence in various locations
224 within continental Asia and the islands of the Sunda Shelf and New Guinea. Third, we
225 use the model to determine which time budget variables are most responsible for
226 limiting the taxon's biogeographic distribution. Fourth, we use the model to determine
227 the possible causes of the historical extinction of gibbon populations in China.
228 Finally, we use the results to assess how plausible it is that female spacing best
229 explains hylobatid monogamy.

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231 **Methods**

232 *Gibbon data*

233 A database was compiled from the gibbon literature to provide quantitative
234 data on key demographic, environmental and ecological variables. We identified 77
235 studies that provide data on at least some of these (see online *Dataset S1*). Of these,
236 59 provide data on mean group size, 41 provide data on group density or biomass, and
237 29 provide data on activity budgets and other behavioural or demographic variables.

238 In respect of activity budgets, we identify five mutually exclusive states (feeding,
239 moving or travel, resting, social interaction – mainly social grooming – and singing)
240 which, in this sample, between them account for ~100% of time.

241 Inevitably, a number of issues arise when comparing data from different
242 studies. One is that studies vary in the definitions they use for individual behaviour
243 categories; a second is that methods of collecting time budget data vary, both in
244 intensity (e.g. some researchers use group scans, others focal individual sampling) and
245 in coverage (the hours of daylight may not be sampled evenly). While these are
246 certainly issues, past experience developing time budget models suggests that the
247 impact of these methodological issues is at best modest (see Dunbar & Shi 2013).
248 More importantly, the main consequence of variations in definition and procedure is
249 that they increase the error variance in parameter estimates; increased error variance
250 will simply make it harder to obtain significant results, and will therefore bias the
251 statistical analysis in favour of the null hypothesis (i.e. no relationship). Ultimately, of
252 course, the real test is whether our models predict what we see on the ground *despite*
253 these methodological flaws. We should be less concerned with standardisation of
254 definitions or methods (though these should always be encouraged) than with whether
255 models based on them correctly predict what we know to be the case.

256 Where these were given by the original field sources, we also extracted data
257 on a number of ecological and behavioural variables: the percentage of leaf in the diet
258 (%Leaf), percentage of fruit in the diet (%Fruit), the density of gibbon groups
259 (groups/ha), gibbon population biomass (kg/km²), mean day journey length (km),
260 mean territory size (ha), density of fig trees (Figdens, indexed as trees/ha) and number
261 of sympatric primate species. Mean day journey length and time spent moving
262 (transformed into hours spent moving per day) were used to calculate travel speed

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(m/hr). Together, these comprise the set of ecological and covariate variables we will use in model-building.

Data for all sites are provided in online *Dataset S1*.

Climatic and geophysical data

Rainfall, altitude and latitude for each study site are those provided by the cited publications, where these are given. Since latitudinal effects should be symmetrical about the equator, we transform all latitudes into absolute latitude. Since individual studies often do not provide all the indices we need, we have sourced all temperature variables (mean annual temperature, TEMP; mean minimum temperature, TEMP_{min}; mean maximum temperature, TEMP_{max}; mean daily temperature variation, TEMP_{var} [difference between mean minimum and mean maximum temperature]; and the standard deviation of mean monthly temperature, T_{moSD} [an index of seasonality]) from either http://www.globalspecies.org/weather_stations/ (which provides climate data for individual wildlife reserves) or <http://en.climate-data.org> (which provides climate data for civic weather stations). Wherever possible, we gave preference to the first. These climatic variables and the two geographical variables (i.e. latitude and altitude, both of which are determinants of climate) constitute the set of climatic variables that we will use as the main independent variables in model-building.

Previous analyses of weather station data for sub-Saharan Africa have demonstrated that only three climatic variables are needed to predict evapo-transpiration (the principal predictor of tropical primary productivity: Rosenzweig 1968; Le Hou  rou & Hoste 1977; Lo Seen Chong, Mougin & Gastellu-Etchegorry 1993). These are mean ambient temperature, total annual rainfall and an index of seasonality (Williamson & Dunbar 1998; see also Hill 1999). As in our previous

models for African primates, the standard deviation of mean monthly temperature (TmoSD) is our main index of seasonality. This also allows us to include any effects due to the fact that, in some contexts, certain times of the year when resource availability is poor or climate especially challenging may impose limits on what animals can do (see, for example, Dunbar & Shi 2013).

Collinearity between the climate variables in the field site dataset is, in general, low and within conventionally acceptable levels, the only exception being that between rainfall and temperature (mainly due to the effect of the cold, dry habitats in southwest China disproportionately influencing what is otherwise a much less clearcut relationship). At best, this relationship explains only 20% of the variance, and as little as 10% if the Chinese habitats are excluded.

Model-building procedure

For the model-building analysis, we excluded time budget data for one study on a heavily logged habitat (Johns 1986) because it had an implausibly low estimate for time spent feeding (8%, >3 standard deviations below the mean for gibbons, and almost a full SD below the next lowest value) and an implausibly high value for time spent resting (62%, 2.5 SDs above the overall mean, and a full SD above the next highest value), two studies (West Garo Hills, NE India: Alfred & Sati 1986, 1990; Ujung Kulon, Indonesia: P. Kappeler 1984a,b) that had very high values for time spent feeding (>60%, 3.1 and 3.9 SDs above the mean for gibbons, and >1 SD above the next nearest value) and unusually low values for resting time, and one (Tanjong Triang: Ellefson 1974) that had an unusually high value for moving (47%) and low value for rest (7%). Such high values for feeding and moving and low values for resting are indicative of gibbons that were not fully habituated.

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313 In developing any model, we face a choice between detail and generality. It is
314 always possible to build a model that takes every conceivable environmental variable
315 into account, and hence is 100% accurate in its fit to the data. But in doing so, we
316 inevitably lose generality: in order to make predictions about where the taxon can
317 live, we will need to know much more about the particular environmental parameters
318 of an individual location (e.g. soil type and acidity, inclination and tree composition,
319 as well as all the larger scale variables like rainfall and temperature). Using general
320 climate as the basis for a model allows us, at the expense of some loss of accuracy, to
321 be more general, since relevant climate values can be mapped geographically on a
322 continental scale from climate models, as well as being projected backwards and
323 forwards in time.

324 For the purposes of building a time budget model, we need to determine
325 taxon-specific equations for just two key variables (feeding and moving time). The
326 other two main components (enforced resting time and social time) use general
327 primate equations given, respectively, by Korstjens et al. (2010) and Lehmann et al.
328 (2007a). For the purposes of the present model, we shall take time devoted to singing
329 as a constant (at the mean observed value of 4.96%).

330 Enforced resting time, as defined by Korstjens et al. (2010), specifies the
331 minimum amount of time that has to be spent resting as a consequence of high
332 ambient temperatures (to avoid thermal overload or excessive heat loss) and for the
333 digestion of leaves (as a function of climatic conditions and the species' dietary
334 physiology). Enforced resting time differs from observed resting time (time spent
335 inactive, as recorded by field observers) in that observed rest time consists of enforced
336 resting time plus uncommitted time ("free rest") (Korstjens et al. 2010).

337 The time during the day that animals have left over after making their habitat-

specific allocations to feeding, moving, enforced rest and singing (subject to any feedback loops in the equation set) gives us the amount of time in the day that could be devoted to social interaction (grooming). Because grooming time correlates with group size (Dunbar 1992; Lehman et al. 2007; Dunbar & Lehmann 2014), we can use this relationship to specify the maximum size of group that the animals could maintain as a coherent social entity at that location. This does not mean the animals *have* to devote all this time to social interaction: it simply sets the upper limit, and hence the upper limit on social group size.

The social time equation reflects the fact that social grooming is the principal bonding mechanism for primate social groups and increases linearly with group size across primates as a whole (Dunbar 1991; Lehmann et al. 2007; Dunbar & Lehmann 2014). As such, it represents the time investment in social interaction that is necessary to maintain a group's cohesion through time so as to prevent it disintegrating (Dunbar 1992a, 2012). It is worth noting that this increase in time devoted to grooming with group size does not mean that animals in large groups groom with more individuals. Typically they do not. Rather, as group size increases, animals invest increasingly in the handful of core grooming partners that act as their primary coalition partners within the group because these act as their primary defence against the stresses of living in a group (Kudo & Dunbar 2001; Dunbar 2012, 2018a).

As in previous models, we first examined all relevant pairwise plots visually to determine whether any factors correlate nonlinearly with the main time budget variables. There were no nonlinear relationships. We then ran backward stepwise linear regression models with the full set of climatic and ecological variables as predictors of each of the two dependent variables (feeding and moving time). The criterion used in selecting the final equation in each case was based on maximising

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the proportion of variance explained, subject to the requirement that the overall equation and all its individual predictor variables were significant.

Because siamang are more than twice as large as other gibbons, siamang feeding time was first corrected to a gibbon-equivalent (i.e. what it would need to do if it was a conventional gibbon) by rescaling it by the ratio of the metabolic body weights of the two taxa (i.e. $5.5^{0.75}/10.75^{0.75}=0.605$). This allows us to use all the data without having to include an additional factor for taxon or body mass. None of the other gibbon species differed significantly in body weight (see *Dataset S1*).

In extracting these equations from the data, we have ignored the possible influence of phylogeny. As in previous analyses (see Dunbar et al. 2009), we do so principally because almost all of the variables we are concerned with are behavioural rather than anatomical, and are likely to be influenced more heavily by local environmental conditions than by biological inheritance, and thus typically have phylogenetic signals that are close to zero (see Kamilar & Cooper 2013). In effect, we treat all the hylobatids as belonging to a single ecological species. Inter-population analyses of just this kind have previously been successfully used to explore aspects of the behaviour of gibbons (song function: Cowlshaw 1992, 1996). It is perhaps important to note that we are not seeking to describe the “typical” behaviour of individual gibbon species: rather, our concern is to understand the overall pattern of behavioural flexibility and the effects of climate on biogeography across the whole taxon. For convenience, we adopt the convention of referring to the entire group as hylobatids, and differentiate between siamang and all other gibbon species only on the basis of body size.

We then use this set of equations to determine maximum group size for any given location. Since all the equations in the present case are linear, we use analytical

methods and solve directly by substitution:

$$N_{\text{Max}} = (100 - (\text{Feed} + \text{Move} + \text{Rest}_{\text{Enf}})) / (\text{grooming equation})$$

This value is the limiting group size (the maximum ecologically tolerable group size, N_{max}).

In order to exhibit the biogeographical implications of the model in graphable form, we present the main results as 3-dimensional surface plots of maximum group size plotted against (a) absolute latitude and altitude and (b) mean annual rainfall and mean annual temperature. Because a relatively large number of environmental and behavioural variables are involved in the basic model, we need to convert all other variables into functions of just the two variables used for each graph. We do this by using backwards stepwise regression modelling to derive equations for these variables from the data given in online *Dataset S1*. The resulting equations are given in Table 2. Of these, latitude, altitude and temperature are well known to be systematically related.

All statistical analyses and graphs were executed in SPSS v.23.

Testing the model

We test the model by evaluating its ability to predict the presence versus absence of gibbons at a range of locations across the Indian subcontinent, the Indo-China peninsula, Indonesia and the Malay Archipelago, New Guinea and mainland China. For these purposes, we selected a number of locations of known altitude and latitude, which had climatic data available in www.en.climatedata.org. The locations and their climatic and geographical data are given in online *Dataset S2*. Since these analyses predict where hylobatids can and cannot survive, they also allow us to make inferences about the biogeographic distribution of the taxon. In most cases, we

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selected major cities for these purposes, since our question is whether or not gibbons could live in the general area, not whether they live at a particular location. These data also allow us to assess whether gibbons could now live in those provinces of China where they were recorded historically by Turvey et al. (2015), and hence determine whether anthropogenic factors or climate change might have been responsible for their extinction. For these purposes we selected one site at random roughly in the center of each of the provinces of modern China.

Results

The model

Table 1 lists the best fit equations for feeding and moving that were generated by the data, as well as the generic primate equations for enforced resting time and social time. The causal relationships involved are summarized in Fig. 1. For reference, alternative “next best” significant equations for feeding and moving time selected by the backwards stepwise regression are given in the *Online Supplementary Material* (Table S1). Both feeding and moving time increase as habitats become climatically more stressful (low rainfall, low temperature, high altitude, high latitude). Although group size has no effect on feeding time, it has a weak positive effect on moving time even despite the extremely limited variation in hylobatid group size (Fig. 2). Since moving time is the main limiting factor for great apes, and severely limits their biogeographic distribution and group size (Lehmann et al. 2007b, 2008a,b), we ran a separate regression model with just latitude and group size as predictors (Eqn. 2a). It is of significance that, despite an order of magnitude difference in the range of group sizes, the slope coefficient for group size in gibbons is very similar to that for the chimpanzee time budget model (3.08 vs 2.59, respectively).

Across the sample of study populations, percentage of fruit in the diet is determined mainly by ambient temperature (Table 2). In contrast, percentage of leaf in diet increases with the level of environmental stress as reflected in rainfall, altitude and temperature variance (Table 2). Although day journey length was not predicted by any of the climatic variables or by fig tree density, territory size is predicted with a very high r^2 by a complex of climatic variables (Table 2), suggesting that territory sizes get larger as environmental stress increases.

The next step is to use the time budget equations to predict maximum ecologically possible (i.e. limiting) group size, N_{max} , for the sampled populations. To do this, we interpolate the time available for social interaction (i.e. that remaining after removing the time predicted for feeding, moving and enforced rest at the site, plus the constant for singing) into Eqn. (5). For this analysis, siamang predicted feeding time is rescaled back to siamang body mass by reversing the transformation used to convert their feeding time to gibbon-equivalents.

Fig. 3(a) plots maximum tolerable group size, N_{max} , against observed mean population group size for all the gibbon and siamang populations in our sample. The horizontal line demarcates a group size of 2.96 individuals, representing the minimum group size for demographic viability (i.e. demographic stability defined by a population growth rate of $r=0$: this requires two surviving offspring over an average gibbon 17-year reproductive lifespan, and would equate to a group with two adults plus $4/17 = 0.12$ births per year, hence an average of ~ 0.96 dependent offspring aged <8 years at any given time, taking 8 to be the age at dispersal and thus the range of individuals who would be observed in a group). (This value assumes minimal mortality in immature gibbons, which seems broadly to be the case; assuming 50% mortality up to puberty, a figure typical of terrestrial baboons, would only increase

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3 463 minimum viable group size to 2 adults + 1.5*0.96 living offspring = 3.44.) The
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5 464 dashed diagonal line marks the line of equivalence (N_{max} is equal to observed mean
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8 465 population size).

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10 466 Three points may be noted. First, time budget models specify that group sizes
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12 467 should not exceed the predicted maximum size (hence all data points in Fig. 3a should
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14 468 lie above the dashed line). In the present case, 80.5% of the 41 populations have a
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16 469 predicted maximum group size that is larger than their observed mean group size.
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18 470 This is broadly encouraging as far as model fit is concerned, but suggests that
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20 471 something else is needed to account for the exceptions. Second, no population has
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22 472 $N_{max} > 15$ individuals. This suggests that hylobatids are under rather greater ecological
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24 473 constraint than many other Old and New World monkeys (and African great apes),
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26 474 where limiting group sizes are typically > 20 . Third, notwithstanding this, it is clear
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28 475 that many (but not all) populations could, on ecological grounds, live in much larger
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30 476 groups than they actually do. Such groups would inevitably be multi-female.

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33 477 One likely reason why some populations might have maximum predicted
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35 478 group sizes below their observed group sizes is the length of the active day. Gibbons
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37 479 are well known to retire early, often several hours before dusk (Raemaekers 1979;
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39 480 Chivers 1984; Palombit 1997; Fei, Zhang, Yuan, Zhang & Fan 2017), a behaviour
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41 481 that is quite unusual for most primates. As a result, the sampled activity budgets are
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43 482 based on an active day that averages 9.3 ± 0.89 hrs (range 8.1-10.6, $N=9$ sites), despite
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45 483 the fact that daylength in tropical habitats is ~ 12 hrs. Progressively extending the
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47 484 length of the active day in units of 5% suggests that an active day that is 15% (84
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49 485 min) longer than the observed mean would be enough to lift all but the three lowest
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51 486 siamang populations above the demographic viability threshold (Fig. 3b). (This would
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53 487 increase maximum N_{max} for the other populations only to ~ 20 .) Unfortunately, data on
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the length of the active day are not available for any of the populations with $N_{max} < 2.96$. However, an extra 84 min would only increase the mean value of day length to 10.7 hrs, virtually identical to the longest observed day length in our dataset (10.6 hrs in a siamang population: *Dataset S1*).

We ran a sensitivity analysis of the model using the alternative equations for feeding and moving time from Table S1. To do this, we altered one of the equations at a time in the original model from Table 1, keeping all the other equations as in the original model. We also ran a model combining the alternative moving time equation with the two most extreme alternative feeding time equations. Predicted maximum group size for the individual populations in the study site database determined using these alternative equations are highly correlated with those determined using the original model (Table S2; Fig. S1). The alternative moving time equation increases the largest maximum group size from ~15 to ~25, but does not substantively change any of the results. This suggests that the Table 1 model is quite robust. Similarly, decreasing the slope coefficient for group size in the moving time equation to 2.59 (the value for the chimpanzee model) increases the largest N_{max} to ~18, while increasing it to 5.0 decreases N_{max} to ~9, but does not alter the broad pattern or the fact that most mean population group sizes are comfortably below N_{max} .

To provide some indication as to how environmental parameters influence hylobatid biogeography, Fig. 4 plots predicted maximum group size as a function of (a) latitude and altitude and (b) rainfall and temperature. To produce these graphs, we have used the equations given in Table 2 to reduce all the climatic variables in the model to the two indices of interest in each case. This can be expected to increase error variance, with a consequential tendency for reversion to the mean, but the results provide us with an indication of how maximum group size is likely to vary across

habitats. The results suggest that hylobatids do best (i.e. are able to sustain larger groups) in cooler, high altitude, high rainfall habitats at low latitudes (i.e. near the equator). When rainfall is less than ~1500mm per annum, or at latitudes above ~20° (in effect, outside the Tropics), hylobatids are unable to maintain minimally viable groups ($N \approx 3$ individuals) unless they are living at altitudes above ~1000m (i.e. under cooler conditions) (see also Turvey et al. 2015). This last prediction is confirmed by the fact that the northern Chinese populations are at significantly higher altitudes than all the other hylobatid populations (mean altitudes: 1750.0 ± 783.8 SD m vs 401.2 ± 409.5 SD m asl, $N=14$ and 50 respectively; $F_{1,62}=76.2$, $p<0.0001$).

Testing the model

To explore the model's ability to predict hylobatid biogeography, we used a sample of locations within the current hylobatid biogeographic range in southeast Asia and a sample of locations on the Indian subcontinent, mainland China and the islands of the Malay archipelago outside the current hylobatid range. In addition, we have included one representative site in each of the Chinese provinces where gibbons were recorded as occurring prior to 1800 AD (see Turvey et al. 2015). Note that, for the latter cases, the climate data are current, not historical, values. In each case, we predict maximum group size using current local climate

Fig. 5 plots the mean and range of maximum group size predicted by the model for each of these sites grouped by geographical location. Predicted maximum group sizes average 6.6 ± 4.0 SD for 18 sites within the current hylobatid range. In contrast, sites in northeast India and Bangladesh to the west of the Brahmaputra (which forms the northwest boundary of the current gibbon range) average 0.3 ± 0.5 SD ($N=7$ sites), as is the case for locations further west in northwest India (mean

1.6±2.2SD, N=7 sites) (see also Fig. S2). This suggests that, even in the absence of the physical barrier of the Brahmaputra river system, the taxon's ability to expand its range westwards has been limited by substantial tracts of land it would have been unable to cross. Had gibbons been able to bypass these two barriers, they would have found very congenial habitats in the southern parts of the subcontinent (e.g. the Nilgiri Hills and adjacent ranges in the Deccan, and in Sri Lanka), where they would have been able to support groups as large as 8.7±5.1SD (N=7 sites). The island habitats across the Wallace Line (specifically, the Celebes and Papua New Guinea) provide equally gibbon-friendly habitats (mean group size 10.3±1.1SD, N=2 sites), but it seems that, as with most other primates (Brandon-Jones 1998), the gibbons were never able to traverse this major sea barrier. Increasing the length of the available day by 15% (as in Fig. 3b) does not change the overall pattern, or make regions such as northern India or central China any more habitable for gibbons.

More puzzling, however, is the fact that central and east Java (including nearby islands like Bali) provide very suitable habitats (mean N_{max} 10.4±1.2SD, N=7 sites) yet have no gibbon populations. Climatically at least, these habitats seem to be at least as suitable as those in west Java where gibbons do occur (mean N_{max} 10.2±1.9SD, N=7 sites). In contrast, gibbons would now find it impossible to survive in either the central and southeastern provinces of China where they were recorded as living prior to 1800 AD (mean 0.0±0.0SD, N=14 sites) or, perhaps less surprisingly, the provinces of northern China where they did not occur historically (mean 1.3±3.6SD, N=7). This contrasts with locations in southwest China (Yunnan province and Hainan Island) that currently do support gibbon populations, for which the model predicts viable group sizes (mean N_{max} = 4.3±1.7SD, N=3 sites).

563 *What limits hylobatid distribution?*

564 An important feature of time budget models is that they allow us to determine
565 which aspect of the animals' biology is the principal constraint on their ability to
566 occupy habitats, and hence what actually limits their biogeographic distribution. In
567 turn, this tells us something about the taxon's risk of extinction under climate change.
568 To explore this, we plotted the predicted time required for feeding, moving and
569 enforced rest for the *Dataset S2* locations (Fig. 6). Neither feeding nor moving differ
570 consistently between regions where hylobatids are present vs not present, suggesting
571 that neither of these is a major constraint (except, in the limit, the effect of group size
572 on moving time at much larger group sizes); in contrast, resting is significantly higher
573 in regions where they are absent and seems to account for the fact that they do not
574 have sufficient time for social interaction to maintain groups of any significant size.

575 It thus seems that the main problem lies in the extent to which climatic
576 variables oblige them to rest. This is confirmed by an analysis of observed versus
577 predicted time budget allocations in the sample study sites. Fig. 7a plots the ratio of
578 observed feeding and resting time divided by the values predicted by the model
579 equations in Table 1 for those populations in the field study dataset (*Dataset S1*) for
580 whom predicted maximum ecologically tolerable group size is either less than or
581 greater than 5 (the upper limit for observed mean population group size). For these
582 purposes, predicted rest is enforced rest time, while observed rest is total rest time.
583 Feeding time is close to that predicted in both cases, as should be the case. In contrast,
584 observed rest time is significantly less than the minimum predicted ('enforced' rest) in
585 populations where N_{max} is less than 5, whereas populations where N_{max} exceeds 5
586 typically have surplus rest time capacity. The strength of this effect is clear from Fig.
587 7b which plots the difference between observed and predicted rest against the

588 difference between observed group size versus predicted N_{max} for the populations.
589 The correlation is highly significant ($r=-0.847$, $p<0.0001$).

591 Discussion

592 The time budget model for hylobatids that we develop here is broadly similar
593 in form to the models developed for 10 other primate genera. It works well for
594 contemporary populations, predicting presence in most cases where gibbons occur and
595 absence where they do not (Fig. 5). In other cases, it is clear that physical barriers
596 (e.g. habitat conditions in northeast India or the Wallace Line to the southeast) have
597 prevented gibbons expanding into habitats where they could do well. It is important to
598 remember that time budget models predict the largest groups that a taxon can maintain
599 at a given site (in effect, the carrying capacity), not necessarily the actual size of
600 group. They simply set the upper limit defined by local environmental conditions.
601 Since, for primates in general, fertility is almost always adversely affected by
602 increasing group size (van Schaik 1982; Dunbar 2018a; Dunbar, MacCarron & Shultz
603 2018a), animals will generally try to minimize group size in any given location to the
604 extent that this is compatible with the constraints imposed by the local predation risk
605 (Dunbar et al. 2009; Dunbar et al. 2018a).

606 The fact that hylobatids commonly live in groups that are smaller than those
607 allowed by the local ecology has two important implications. First, it implies that,
608 while the model certainly tells us that gibbons are under greater ecological constraint
609 than most Old World monkeys and African great apes, the level of ecological stress is
610 not sufficiently high to force them to live in groups as small as those they actually live
611 in. While it is true that some populations have time budgets with little or no spare
612 capacity, many do not (Fig. 7). This conclusion is supported by the fact that, unlike

most other monkeys and apes, hylobatids commonly go to their night rest mid-afternoon (Raemaekers 1979; Srikosamatara 1984; Palombit 1999; Fei et al. 2017), implying that they are not under significant time pressure. Second, it suggests that predation risk must be low for hylobatids; group size evidently does not provide gibbons with a significant anti-predator advantage, so they can afford to minimize group size in order to maximize fertility (see Dunbar et al. 2009; Bettridge, Lehmann & Dunbar 2010; Dunbar, MacCarron & Robertson 2018b). Indeed, in comparison to almost all other primates, cases of predation are conspicuous by their absence from the gibbon literature (Reichard 2003).

This calls into question the claim that gibbons are monogamous because females are forced by their ecology to forage on their own rather than in groups (van Schaik & van Hooff 1983; Rutberg 1985; Komers & Brotherton 1997; Brotherton & Komers 2003; Lukas & Clutton-Brock 2013). Contrary to this claim, it seems that most populations could in fact maintain larger groups (range 5-12 animals), especially in the more southerly parts of the Malay archipelago. Groups of this size could easily include 2-4 reproductive females (across all primates, reproductive females form a very consistent 30-35% of the group: Dunbar et al. 2018a,b). Even if females lived alone, it would be possible to have more of them within a gibbon group's current territory, and males would have access to more than one reproductive female if they pursued a roving male strategy (much as orang utan males do) (see also van Schaik & Dunbar 1990; Bartlett 2009). This implies, as suggested by van Schaik & Dunbar (1990) and Opie, Atkinson, Dunbar & Shultz (2014), that monogamy must have evolved in response to factors other than ecological constraints.

One of the main benefits of the time budget model approach is that it provides insights into the behavioural and physiological constraints that limit a taxon's ability

638 to successfully occupy different kinds of habitats. As with previous primate time
639 budget models, feeding time is strongly influenced by predictors of habitat quality: in
640 the hylobatid case, the main determinants of feeding time (rainfall, altitude and
641 latitude, with the latter two both being core predictors of temperature) are all variables
642 that influence both tree species composition and the nutritional quality of vegetation
643 (see also Marshall & Leighton 2006). As food quality falls, animals will need to spend
644 longer feeding to meet their nutritional requirements.

645 In contrast, time spent moving remains largely independent of the climatic
646 variables, being affected only by absolute latitude and group size. Models for African
647 genera have also noted that moving time tends to be independent of environmental
648 variables and is often close to being a constant (Dunbar et al. 2009), mainly because,
649 in order to save time for other more pressing activities, animals respond by increasing
650 travel speed rather than increasing travel time when they need to travel further
651 (baboons: Dunbar 1992a). There is some suggestion that gibbons also do this: travel
652 speed increases as climatic conditions deteriorate (i.e. when temperatures are high and
653 latitude is low, although the effect is weak and not significant (Fig. S3). However,
654 since there is inevitably a limit on the speed at which animals can travel, at some point
655 animals simply have to devote more time to moving. The substantive issue for day
656 journey length (and hence moving time) is, at least for non-folivores (Snaith &
657 Chapman 2007), group size: the area that has to be covered to allow every animal in
658 the group to meet its nutritional requirements is inevitably a linear function of the
659 number of animals in the group, forcing the group to travel further each day (Dunbar
660 et al. 2009). The great apes are especially susceptible to the effects of group size on
661 moving time, and it is this that is mainly responsible for chimpanzees' fission-fusion
662 form of sociality (Lehmann et al. 2007b, 2008a,b). It is significant that, despite the

663 very limited variance in the size of their groups, the group size slope parameter for the
664 gibbons is similar to that for chimpanzees. For groups as small as those found in
665 gibbons, the impact of this effect will, of course, be modest; but it does mean that
666 when groups are larger than ~10 the impact will rapidly becomes prohibitive (as is the
667 case for African great apes: Lehmann et al. 2007b [Fig. 7]).

668 However, it seems to be enforced resting time that is the main constraint for
669 hylobatids (Figs. 6 and 7). Since this is mainly affected by ambient temperature, it
670 might explain why social groups get larger in northern populations irrespective of
671 species (Fig. 8). Ultimately, however, the northern extension of hylobatids is
672 constrained by the fact that feeding and moving time are also positive functions of
673 latitude (and hence, in effect, declining temperatures). A comparison of the slope
674 parameters for latitude in Tables 1 and 2 indicates that the additive effects of the
675 latitude coefficients for feeding and moving time increase at a combined rate that is
676 ~4 times the savings in terms of the slope for enforced rest ($0.363 + 0.462 = 0.825$ vs
677 $1.33 \times 0.158 = 0.210$), so that time budgets become progressively squeezed as animals
678 occupy increasingly high latitude habitats. At the latitude of Mt Wuliang in Yunnan
679 Province, latitude has added a net 15 percentage points to the time budget.

680 There is a widespread perception that *Nomascus* populations live in larger
681 groups than all other hylobatids, hence the fact that they have polygamous groups. Up
682 to a point, this is true; however, this is not true for all *Nomascus* populations (Fig. 8).
683 Moreover, *Hoolock* populations also have large groups at high latitudes without these
684 being polygamous. Mean population group size is in fact a cubic function of absolute
685 latitude (Fig. 8). Group size does increase steadily up to around 20°N (roughly the
686 latitude of Chaing Mai in northern Thailand), but then it levels off only to drop
687 precipitately after latitude 25°N (roughly the latitude of Mt Wuliang in southwestern

China). This decline at very high latitudes is suggestive of populations living at the limits of their range, and indeed mean group size at these highest latitude populations is only just above the minimum for demographic viability.

It is evident from Fig. 3(b) that siamang incur an additional cost due to the fact that their larger body size imposes a higher feeding time demand. A convincing explanation for their larger body size remains elusive, especially given the fact that their range overlaps that of other gibbons and the two taxa are often sympatric. As a result, however, they are often under significant time budget pressure. There is some evidence to suggest that they make time budgeting adjustments that might be sufficient to reduce this pressure: for the limited sample available, it seems that siamang devote less time each day to both singing and social interaction than gibbons do, and they have a longer active day (Table 3). For this limited sample, the net gain for siamang is equivalent to increasing the length of the active day by ~17% (the combined effect of a longer active day adjusted for less time spent socialising and singing) while only having to increase the actual length of the active day by ~10%. That is sufficient to just lift the lower siamang populations in Fig. 3(b) into the minimum viability zone. Nonetheless, siamang populations are clearly under more ecological pressure than is typical of the other gibbons (Fig. 3).

Gibbons share with the great apes a dietary physiology specialized for frugivory; indeed, their feeding time equation is very similar to that of the gorilla, though their size precludes their being able to survive on low quality folivorous fallback foods in the way gorillas can. Although the larger-bodied siamang exhibit some capacity in this direction (Raemaekers 1979; Palombit 1995), their ability to resort to a heavily folivorous diet is likely to be considerably less than the gorilla's simply because of their smaller body size. Most years aside, fruits invariably have a

713 more patchy distribution than leaf, and the travel demands imposed by this may be
714 part of the problem that affects gibbon time budgets, and hence limits group sizes.

715 Gibbon ecology appears to be quite tightly defined: their geographical range is
716 surprisingly limited. They are currently confined to the Indochina peninsula (bridging
717 out into southwest China) and the associated islands of the Malay archipelago
718 (Sumatra, Java and Borneo). They appear to have been prevented from extending
719 southeastwards by the Wallace Line (despite the fact that the islands beyond would be
720 perfectly habitable for them: Fig. 5), while their capacity to encroach into the Indian
721 subcontinent has been limited by the climate west of the Brahmaputra (Fig. 6).
722 Nonetheless, within their core distribution they seem to be under much less ecological
723 constraint than we might have anticipated given the female-dispersion explanation for
724 their adoption of a monogamous mating/social system.

725 Historically, gibbons are known to have occurred at much higher latitudes in
726 China than they do now. The model suggests that current climate in these areas makes
727 these habitats completely unsuitable for gibbons. Although the historical extinction of
728 these northern populations has been attributed to anthropogenic factors (Fan 2017), in
729 fact these habitats would not support gibbons now even if there were no humans
730 living there. There are only three plausible explanations for the disappearance of these
731 populations. One is that the populations in question had evolved novel adaptations to
732 these environments, such that the slope parameters on their feeding and/or moving
733 equations were radically different from those for gibbons from further south, but that
734 humans caused their extermination nonetheless. However, to suggest that these
735 northern gibbons were doing something completely different to all other New and Old
736 World anthropoid primates as well as other gibbons implies that they could not have
737 been conventional primates, and that should be, at best, an explanation of last resort.

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3 738 A second possibility is that humans had released animals in these areas in order to
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5 739 have them as exotics (something that humans seem to be especially prone to do), even
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7 740 though the habitats were not really suitable for them; as a result, the populations went
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9 741 extinct once they were no longer being replenished (a situation not too dissimilar to
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11 742 that of *Macaca sylvanus* on Gibraltar). This is a possibility, but the fact that gibbons
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13 743 seem to be unable to survive there at all would imply the need for constant
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15 744 replenishment, and this seems unlikely over such a wide area and such a long time
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17 745 period. The third option is that the climate has changed in the past 300 years in ways
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19 746 that would have led to the inevitable extinction of these populations (with or without
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21 747 the assistance of humans).

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26 748 A potential culprit in the latter respect is the Little Ice Age of 1645-1715,
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28 749 which resulted in average world temperatures falling by $\sim 1^{\circ}\text{C}$. In China, this resulted
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30 750 in a significant increase in climatic seasonality. The cultivation of Mediterranean-type
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32 751 citrus crops was abandoned in Jiangxi Province (one of the provinces that historically
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34 752 harboured gibbons: Turvey et al. 2015) after the 1750s, despite their having been
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36 753 cultivated there for many centuries (Reiter 2000). Guangdong (another of the
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38 754 provinces where gibbons have been documented) experienced a particularly cold, dry
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40 755 spell, accompanied by an unusually high frequency of typhoon strikes, after the Little
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42 756 Ice Age (Liu, Shen & Louie 2001). Ameca y Juárez, Mace, Cowlshaw, Cornforth &
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44 757 Pettorelli (2013) have shown, for mammals as a whole and primates in particular, that
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46 758 high frequencies of cyclones and droughts correlate with elevated extinction risk
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48 759 (indexed by the number of taxa classified as “threatened”), with the southeast corner
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50 760 of China being especially prone to this effect. Turvey et al. (2015) noted that there
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52 761 was increasing fragmentation of these populations from 1700 onwards (with a marked
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54 762 upturn in fragmentation from around 1900), with a 50-100 year lag to last reported
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occurrence. This is indicative of populations struggling to survive.

A comparison of the distribution of climatic variables in southwest China (where gibbons live now) and central and southeast China (where they lived historically) suggests that the only climate variables in which these two regions differ significantly are $TEMP_{moSD}$ (standard deviation of mean monthly temperature across the year, an index of seasonality) and annual rainfall (Table 4). (We do not consider latitude or altitude since these cannot have changed historically.) This may well reflect shifting patterns in the latitudinal distribution of the monsoon circulation, allowing an extension of a more seasonal monsoon climate further into mainland China after the Little Ice Age. A 65% increase in rainfall and a dramatic reduction in seasonality would be required to allow gibbons to survive in the southeastern provinces as well as they currently do in the southwest.

The bottom line seems to be that even if anthropogenic factors have been important in the final demise of these populations (as both Turvey et al. 2015 and Fan 2017 have suggested), these populations' sensitivity to anthropogenic factors is likely to have been exacerbated by environmental effects on the animals' increasing inability to cope as climate changed. The results in Fig. 4, for example, imply that, historically, declining populations at high latitudes are likely to have become locked into mountain-top retreats as climate deteriorated, leading to small, isolated pockets that are inevitably more vulnerable to extinction in the face of environmental shocks (Dunbar 1998; Cowlishaw & Dunbar 2000).

One final puzzle is the fact that gibbons are not found east of the Dieng Mountains in central Java. Kappeler (1984a,b) attributed this to the kinds of forests found in the eastern half of the island, which tend to be more deciduous. Fossil gibbons have been recorded at Trinil, central Java (in the Lower/Middle Pleistocene

deposits that produced the hominins: Ingicco, Vos & Hoffman 2014) and at the Ngandong and Gunung Dawung hominin sites in east Java (Storm & Vos 2006) (see also Jablonski & Chapin 2009), so this area clearly did once support gibbon populations. The fact that west Java was one of the refugia for gibbons during the dry phases of the late Pleistocene Ice Ages (Brandon-Jones 1998) makes this all the more puzzling. It may be that anthropogenic factors resulted in their demise, although these are unlikely to have kicked in until historical times.

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Legends to Figures

Fig. 1. Flow chart for the gibbon time budget model, summarising the causal relationships between climatic and behavioural variables based on the equations given in Table 1. Solid lines: positive effects; dashed lines: negative effects.

Fig. 2. Time spent moving plotted against mean population group size. Dashed line is least squares regression line. Filled symbols: gibbons; unfilled symbols: siamang. Source: online *Dataset S1*.

Fig. 3. (a) Maximum ecologically tolerable group size predicted by the time budget model, plotted against observed mean group size for different gibbon (filled symbols) and siamang (unfilled symbols) populations. (b) Predicted maximum group size allowing for a 15% increase in the length of the active day. The solid line demarcates the minimum group size for demographic viability (2 adults plus 0.96 immatures: see text for details); dashed line in (a) is the line of equilibrium (N_{max} = observed mean).

Fig. 4. Maximum group size predicted by the time budget model for different combinations of (a) latitude and altitude and (b) annual rainfall and mean temperature.

Fig. 5. Mean ($\pm 95\%$ CI) maximum group size (N_{max}) predicted by the time budget model for habitats within different geographic regions. Filled circles: regions

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3 1089 where gibbons currently live; unfilled circles: regions where gibbons do not
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5 1090 currently live; grey circle: gibbons historically present before 1800 AD. Dashed
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7 1091 line: minimum group size for demographic viability (2.96 individuals: see text).
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10 1092 Present: 18 sites within the current biogeographic distribution of gibbons within
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12 1093 Indo-China, Sumatra and Borneo. W Java: 3 sites in west Java where gibbons
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14 1094 occur; E Java: 8 sites in Java east of the Dieng Mountains where gibbons do not
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16 1095 currently occur; E Indonesia: 2 sites on Celebes and western Papua New Guinea,
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18 1096 east of the Wallace Line where gibbons do not occur; NW and NE India: 8 and 7
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20 1097 sites, respectively, in northwest and northeast India (including Bangladesh) where
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22 1098 gibbons do not occur; S India: 7 sites in southern India plus Sri Lanka; SW
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24 1099 China: 3 sites within current gibbon range in Yunnan and Hainan. SE China: one
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26 1100 site from each province in southern China where gibbons were recorded as being
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28 1101 present in historical documents before 1800 AD (N=14 site). N China: one site
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30 1102 from each province in northern China (including Tibet) where gibbons were not
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32 1103 recorded historically (N=7 sites). Locations and climate data from online *Dataset*
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34 1104 *S2*.
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42 1106 Fig. 6. Mean ($\pm 95\%$ CI) (a) feeding time, (b) moving time and (c) enforced rest time
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44 1107 predicted by the time budget model for the regional sites shown in Fig. 5. Moving
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46 1108 time does not include an adjustment for group size. Regions and symbols as for
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48 1109 Fig. 5.
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51 1110
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53 1111 Fig. 7. (a) Mean ($\pm 95\%$ CI) ratio of observed to predicted time spent feeding and
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55 1112 resting for populations for which predicted $N_{max} < 5$ (unfilled symbols) or N_{max}
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57 1113 > 5 (filled symbols). Predicted rest time is enforced rest time. (b) Difference in
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1114 rest time (observed minus predicted by model) for individual hylobatid
1115 populations plotted against difference in group size (observed minus *Nmax*).

1116 Source: Dataset S1.

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1118 Fig. 8. Mean observed group size for individual populations plotted against absolute

1119 latitude for the population. The best fit equation (dashed line) is Group size =

1120 $3.71 - 0.014 * \text{Lat} + 0.012 * \text{Lat}^2 - 0.0004 * \text{Lat}^3$ ($r^2=0.159$, $F_{3,50}=3.15$, $p=0.033$).

1121 Source: Dataset S1.

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Table 1. Regression equations for the hylobatid model.

Variable	Equation	r ²	F	df	p
(1) Feed time (%)	41.634 – 0.003*RAIN – 0.009*ALT + 0.363* LAT _{abs} ^Ω	0.501	5.03	3,15	0.013
(2) Moving time (%)	17.316 + 0.462*LAT _{abs}	0.166	3.59	1,18	0.074
(2a) Moving time (%)	5.297 + 0.290*LAT _{abs} + 3.080*N	0.220	1.98	2,14	0.175
(3) Rest _{ENF} time (%)¶	-23.24 + 1.33*TEMP + 0.259*LEAF + 6.12*T _{moSD}	from Korstjens et al. (2010)			
(4) Social(%)†	2.968 + 0.109N	0.440	29.90	1,38	<0.0001
(5) Time budget	100 = FEED + MOVE + REST _{ENF} + SING + SOCIAL				
(5a)	100 = FEED + (5.297 + 0.290*LAT _{abs} + 3.080*N) + (REST _{ENF} + REST _{FREE}) + SING + (2.968 + 0.109*N)				
(5b) Nmax§	= ((100 – FEED – (5.297 + 0.290*LAT _{abs})– REST – 4.96‡) + 2.968)/(0.109 + 3.080)				by substitution

Ω Absolute latitude

¶ Rest time consists of two components: enforced rest (REST_{ENF}, imposed on the animal by climatic conditions and its dietary strategy) and free rest (REST_{FREE}) that represents uncommitted time that can be allocated to any other activity when required (see Korstjens et al. 2010).

† Recalculated from Lehmann et al (2007a)

§ Maximum ecologically tolerable group size (the maximum group size that will allow the time budget to be balanced)

‡ 4.96 = average percentage of day devoted to singing by gibbons

Table 2. Supplementary equations

Variable	Equation	r ²	F	df	p
TmoSD **	10.440 + 0.093*LAT _{Abs} - 0.001*ALT - 0.821*TEMP + 0.588*TEMP _{min}	0.597	17.75	4,48	<0.0001
Mean temperature (°C)	27.651 - 0.003*ALT - 0.158*LAT _{Abs}	0.672	60.33	2,59	0.0001
Altitude (m)	4145.354 - 147.236*TEMP + 0.023*RAIN	0.575	37.14	2,55	<0.0001
Temp _{var} †	10.196				
Absolute Latitude	48.732 - 0.004*RAIN - 1.142*TEMP	0.580	35.23	2,51	<0.0001
Rain	3279.523 - 53.609*LAT _{Abs} + 0.025*ALT	0.286	10.60	2,52	<0.0001
Temp _{min} (°C)	-5.776 + 1.033*TEMP	0.922	623.63	1,53	<0.0001
Territory size (ha)	2498.88 + 0.077*ALT - 477.830*TEMP _{min} + 401.718*TEMP _{max} - 477.03*TEMP _{var} - 32.260*TEMP _{moSD} - 0.032*RAIN	0.894	19.60	6,14	<0.0001
Leaf in diet (%)	99.430 - 1.000*LAT _{abs} + 0.009*ALT + 7.012*TEMP _{var} + 7.879*TEMP _{min} - 9.779*TEMP _{max}	0.611	5.04	5,16	0.006
Fruit in diet (%)	-51.27 + 10.65*TEMP - 7.71*TEMP _{min}	0.382	6.19	2,20	0.008
Active day (hr) ^Ω	11.273 - 0.001*RAIN	0.905	19.15	1,2	0.048

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** Standard deviation of monthly mean temperatures across the year
† Difference between mean monthly maximum and mean monthly minimum temperature
Ω Gibbons only

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Table 3. Time budget differences between siamang and gibbons

Variable	Siamang		Gibbons*	
	Mean±SD	N§	Mean±SD	N§
% fruit in diet	40.7±16.1	6	57.9±18.7	22
% leaf in diet	34.6±11.6	4	33.7±30.4	22
Feed (%)	49.6±6.8	5	34.4±8.5	19
Move (%)	18.0±5.8	5	21.4±7.9	16
Rest (%)	34.4±10.1	5	34.7±9.8	18
Sing (%)	2.0	1	4.96±3.1	10
Social time (%)	3.0	1	5.87±3.5	12
Active day (hr)	10.1±0.3	2	9.06±0.9	7
Net difference in available time†		16.8%		

* sites used in time budget analyses only

§ Number of studies

† Equivalent change in gross time budget achieved by the siamang compared to gibbons (~3% by reducing singing, ~3% by reducing social time and ~11% by increasing length of active day)

Table 4. Comparison of climatic variables between southwest China (where gibbons currently exist) and southeast China (where they were present historically, but no longer are).

Provinces of China	Gibbons present	TEMP	TEMP _{min}	TEMP _{max}	TEMP _{var}	TEMP _{moSD}	RAIN
Southwest	current	20.6	16.5	24.6	8.1	3.8	1973
Southeast	historical	16.6	12.5	20.9	9.7	8.1	1163
	F _{1,15}	2.50	1.88	2.61	0.21	19.97	9.22
	p	0.135	0.191	0.127	0.657	0.0005	0.008

Environment and Time as Constraints on the Biogeographical Distribution of Gibbons

**R.I.M. Dunbar, Susan M. Cheyne, Daoying Lan, Amanda Korstjens,
Julia Lehmann & G. Cowlshaw**

Online Supplementary Material

Alternative model equations

Table S1 provides alternative multivariate equations for feeding and moving time generated by the backwards regression model. Most of the slope coefficients are similar across the alternative equations. However, note that the magnitude of the effect of group size on moving time for Equation (2c) is considerably less. Nonetheless, the values for both Equations (2a) and (2c) straddle the observed value for the chimpanzee model.

Table S2 gives the correlation values for maximum ecologically tolerable group size predicted by the original model based on Table 1 in the main text and the various alternative feeding and moving time equations given in Table S1. All alternative statistical models yield predicted maximum group sizes that correlate significantly with those predicted by the original model based on the equations of Table 1.

Fig. S1 plots predicted maximum group size for individual study sites in the main dataset (online *Dataset S1*) against the equivalent value predicted by substituting the various alternative feeding and moving time equations. Alternative feeding time equations do not have a significant effect on predicted maximum group size. Note that the alternative moving time equation predicts slightly larger group sizes than those predicted by the original model of Table 1 because the effect of group size on moving time in Equation 2c is lower (1.72 vs 3.08), yielding slightly larger group sizes. Nonetheless, the difference in the size of the largest groups predicted is modest (25 vs 15), the outcomes do not differ significantly (Table S2). Hence, using these alternative equations does not change any of the main conclusions.

Table S1. Alternative regression equations for feeding and moving time for the hylobatid model.

Variable	Equation	r ²	F	df	p
(1) Feed time (%)	41.634 + 0.363* LAT _{abs} – 0.009*ALT – 0.003*RAIN	0.501	5.03	3,15	0.013
(1a)	53.955 + 0.243*LAT – 0.010*ALT – 0.472*TEMP + 0.243*TEMPVAR - 0.004*RAIN	0.340	2.86	5,13	0.059
(1b)	55.879 + 0.279*LAT _{abs} – 0.010*ALT -0.485*TEMP -.004*RAIN	0.520	3.80	4,14	0.027
(1c)	33.230 + 0.469*LAT _{abs} – 0.009*ALT	0.461	6.85	2,16	0.007
(2) Moving time (%)	17.316 + 0.462*LAT _{abs}	0.273	5.25	1,14	0.038
(2a)	5.297 + 0.290*LAT _{abs} + 3.080*N	0.220	1.98	2,14	0.175
(2b)	29.196 + 0.306*LAT – 0.365*TEMP _{max}	0.294	2.71	2,13	0.104
(2c)	19.409 + 0.303*LAT _{abs} – 0.275*TEMP _{max} + 1.718*N	0.312	1.81	3,12	0.199

Table S2. Correlations between maximum group size predicted by the original model of Table 1 and a model using the different equations for feeding and moving time given in Table S1, for the actual hylobatid study sites (as listed in online *Dataset S1*). In each case, one equation is substituted, with all the remaining equations as in the original model of Table 1.



Pearson Correlations

		Feed Eqn 1a	Feed Eqn 1b	Feed Eqn 1c	Move Eqn 2c	Feed Eqn 1a with Move Eqn 2c	Feed Eqn 1c with Move Eqn 2c
Original model	Correlation	0.994	0.995	0.988	0.996	0.987	0.986
	p (2-tailed)	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001
	N	51	51	51	51	51	51
Feed Eqn 1a	Correlation		0.999	0.974	0.996	0.997	0.978
	p (2-tailed)		<0.0001	<0.0001	<0.0001	<0.0001	<0.0001
	N		51	51	51	51	51
Feed Eqn 1b	Correlation			0.975	0.997	0.997	0.980
	p (2-tailed)			<0.0001	<0.0001	<0.0001	<0.0001
	N			51	51	51	51
Feed Eqn 1c	Correlation				0.984	0.966	0.997
	p (2-tailed)				<0.0001	<0.0001	<0.0001
	N				51	51	57

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Move Eqn 2c	Correlation					0.995	0.989
	p (2-tailed)					<0.0001	<0.0001
	N					51	51
Feed Eqn 1a with Move Eqn 2c	Correlation						0.977
	p (2-tailed)						<0.0001
	N						51

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Figure S1

Maximum group size predicted by the original model using the Table 1 equations plotted against maximum group size for all populations in the sample of studies (*Dataset S1*), using the alternative Feed and Move equations given in Table S1. Except for the last pair of graphs, each alternative equation is substituted individually, and all other equations are held constant as in the original model of Table 1. Filled symbols: gibbon populations; unfilled symbols: siamang populations. Dashed line: line of equivalence (the predicted maximum group size is the same for both models).

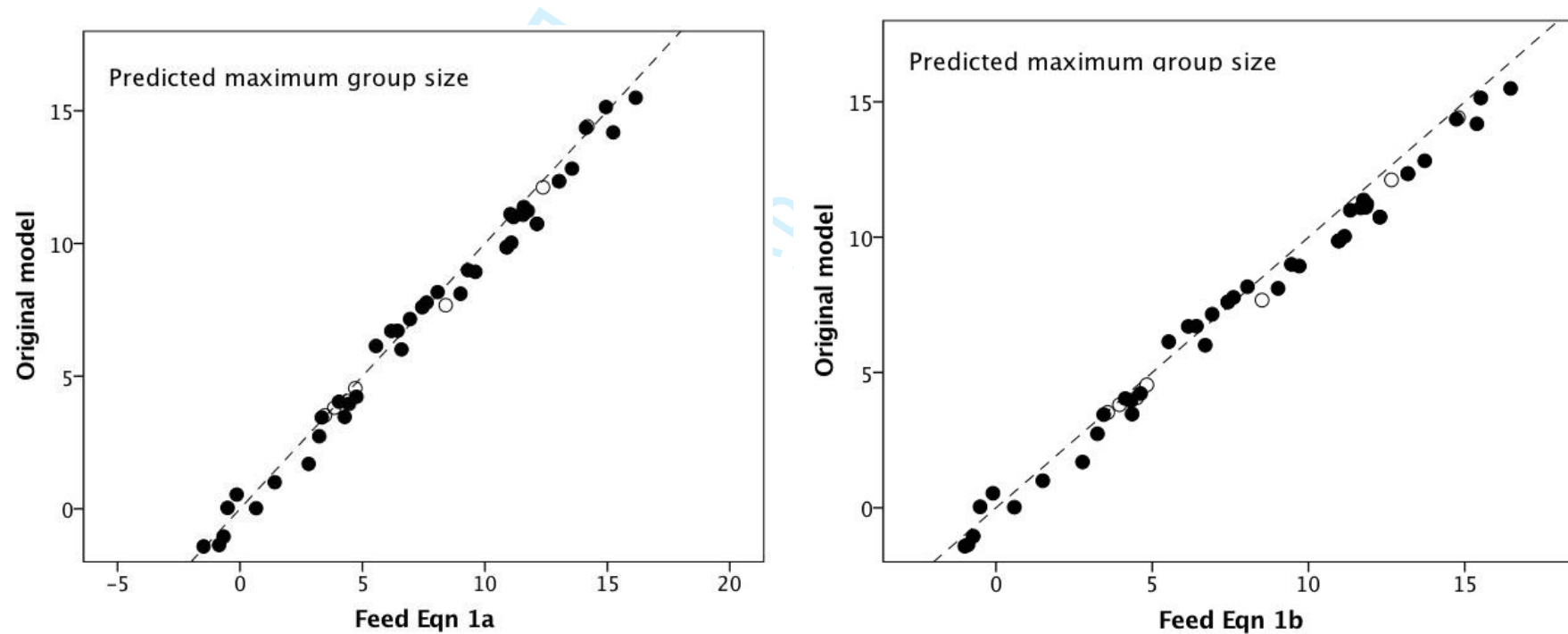


Fig. S1 (contd)

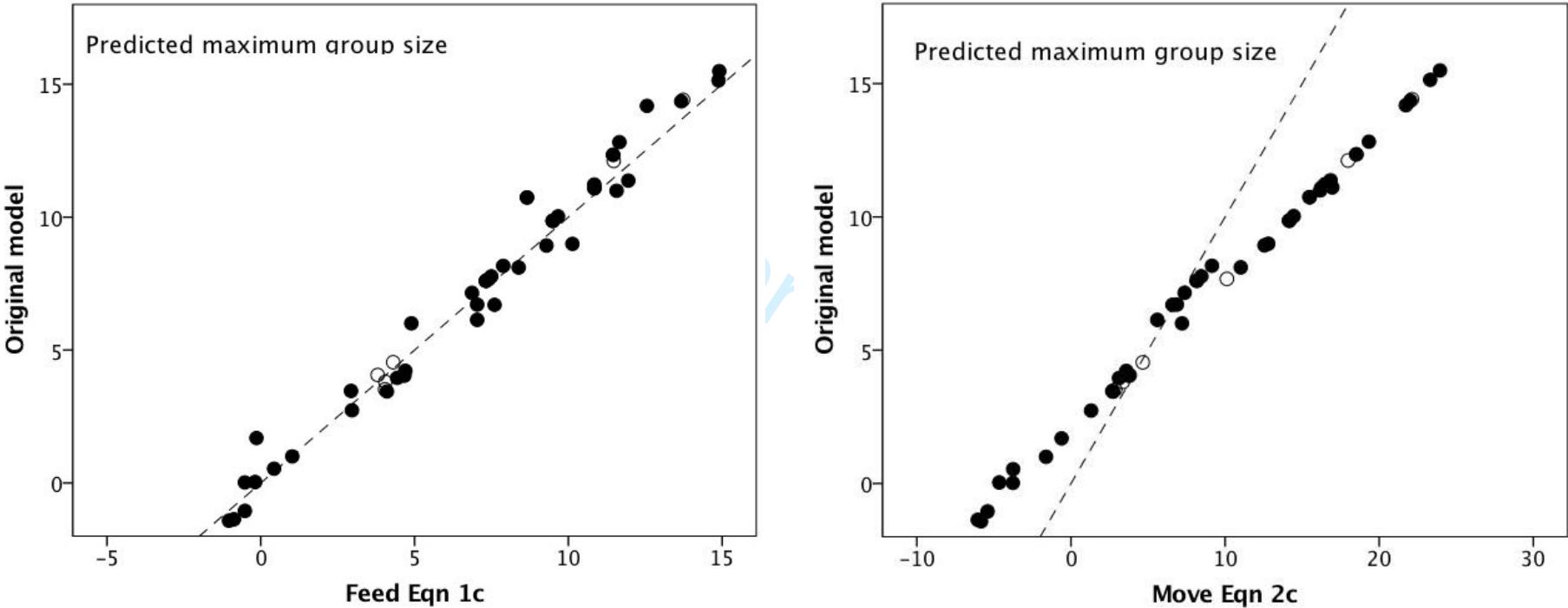
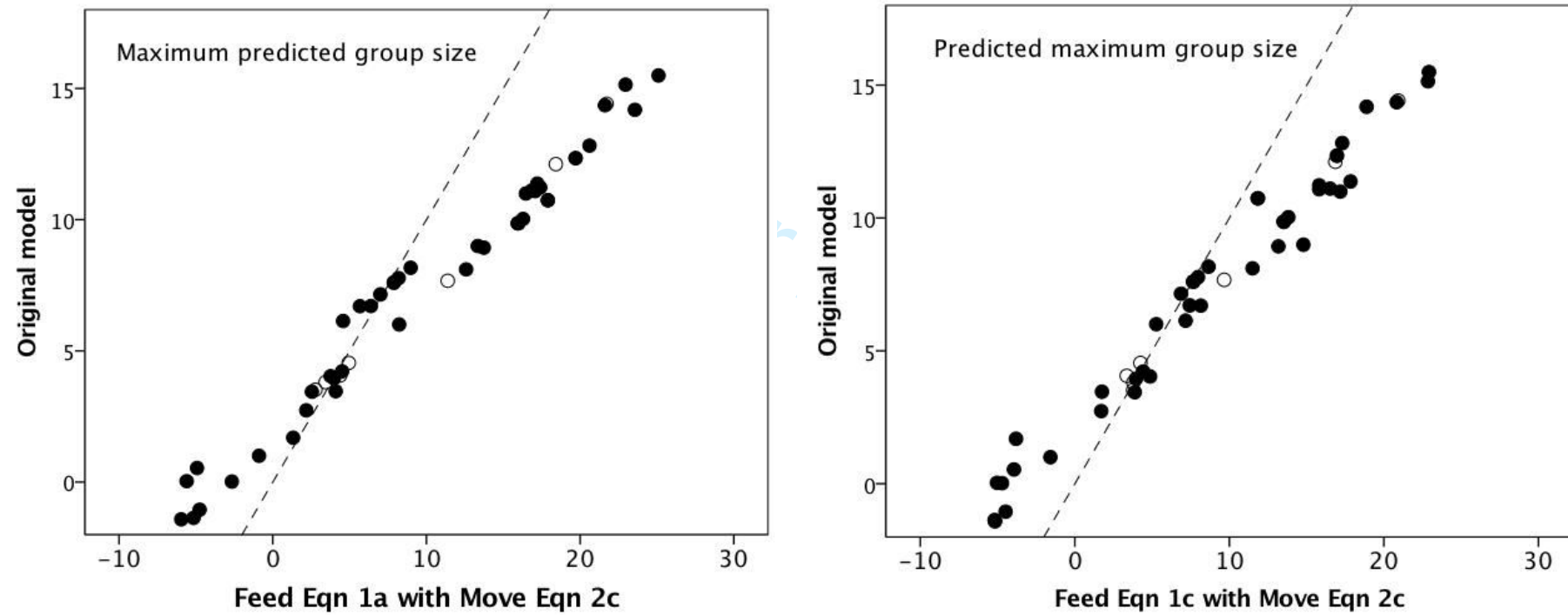


Fig. S1 (contd)



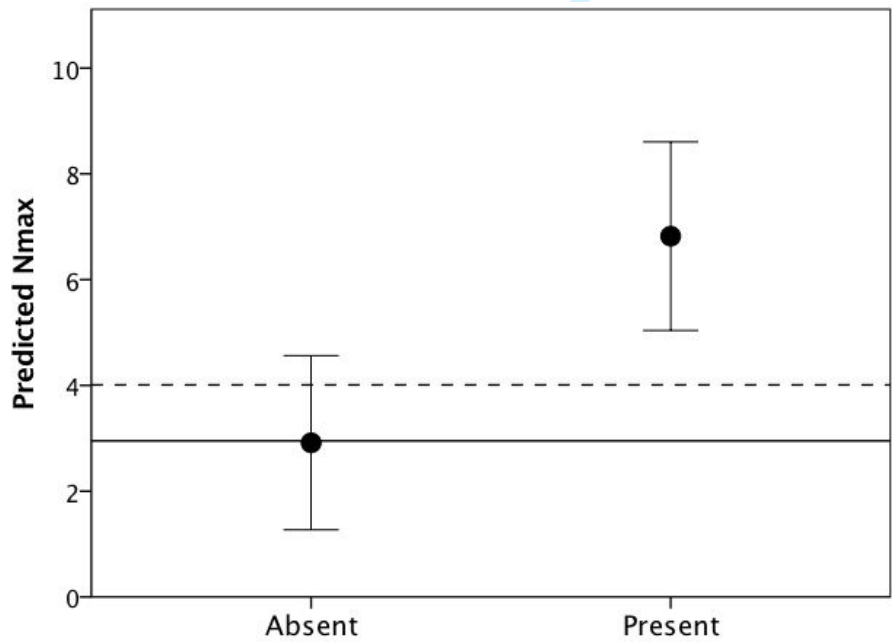
Model predictions

Fig. S2 plots the mean ($\pm 95\%$ CI) predicted maximum group size (N_{max}) for sites in countries where hylobatids are known to live and those in adjacent geographical regions (which should be accessible to them) where they do not. These values are calculated from the climate data in *Dataset S2* for those regions listed as within their current geographic range, plus SE China (essentially the provinces where gibbons occurred prior to 1800), NE India (the area north and west of the Brahmaputra in Bangladesh and northeastern India), and central and eastern Java. Indonesia east of the Wallace Line is excluded.

The difference is highly significant (means 2.92 vs 6.8, $N=31$ and 23 , respectively; $F_{1,52}=10.71$, $p=0.002$). Note that, in areas where gibbons do not live, half the N_{max} values are below the group size required for demographic viability (2.96 individuals), and most are below the observed mean size of 4.01 individuals. In contrast, all values for locations where gibbons live are above the observed mean group size.

Figure S2

Mean ($\pm 95\%$ CI) predicted maximum group size (N_{max}) for sites in countries where hylobatids are currently found, compared to that for geographically adjacent areas where they do not now occur (see main text for details). The solid line is the calculated minimum group size for demographic viability (2.96 individuals of all ages: see main text); the dashed line is the observed mean group size across all gibbon populations. Source: *Dataset S2*



Speed of travel

We calculate average speed of travel across the day as day journey length (in metres) divided by the number of hours in the day (12) multiplied by the proportion of the day devoted to moving. Speed of travel is weakly predicted by latitude, altitude and temperature:

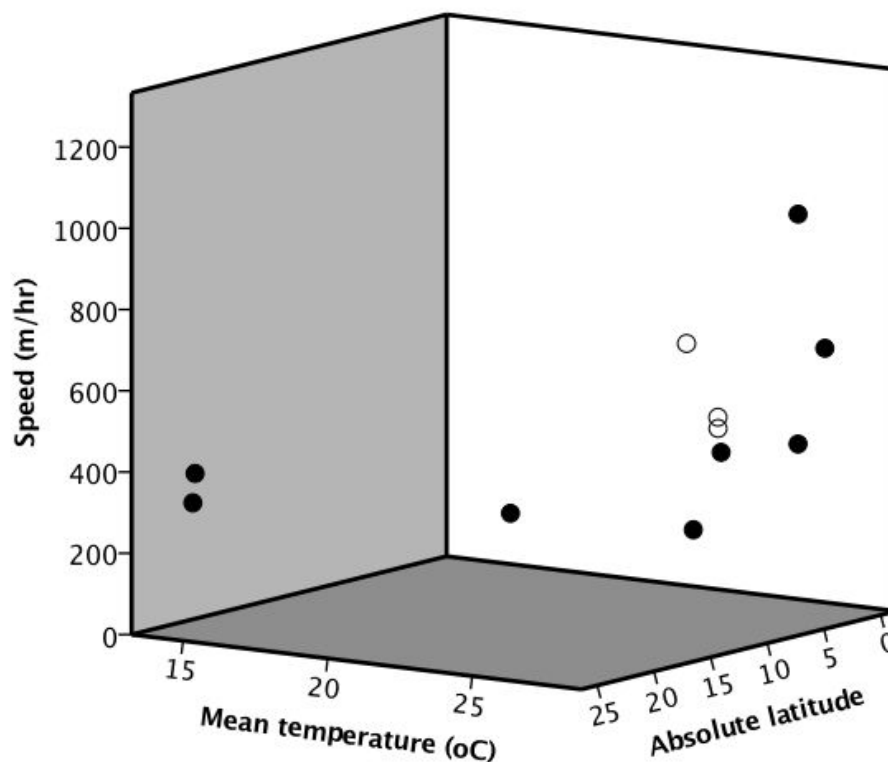
$$\text{SPEED (m/hr)} = -621.2 - 25.7 * \text{LAT}_{\text{Abs}} + 0.4 * \text{ALT} + 46.5 * \text{TEMP}$$

($r^2=0.674$, $F_{3,7}=1.94$, $p=0.211$).

Fig. S3(a) plots speed against absolute latitude and mean temperature. There is a very slight suggestion that the relationship might be quadratic (U-shaped) in temperature, with speed increasing when ambient temperatures are both low and high, with a minimum at temperatures around 20°C. Alternatively, the relationship may be better described by a power curve in which speed increases dramatically above ~22°C at latitudes <5° either side of the equator.

Figure S3(a)

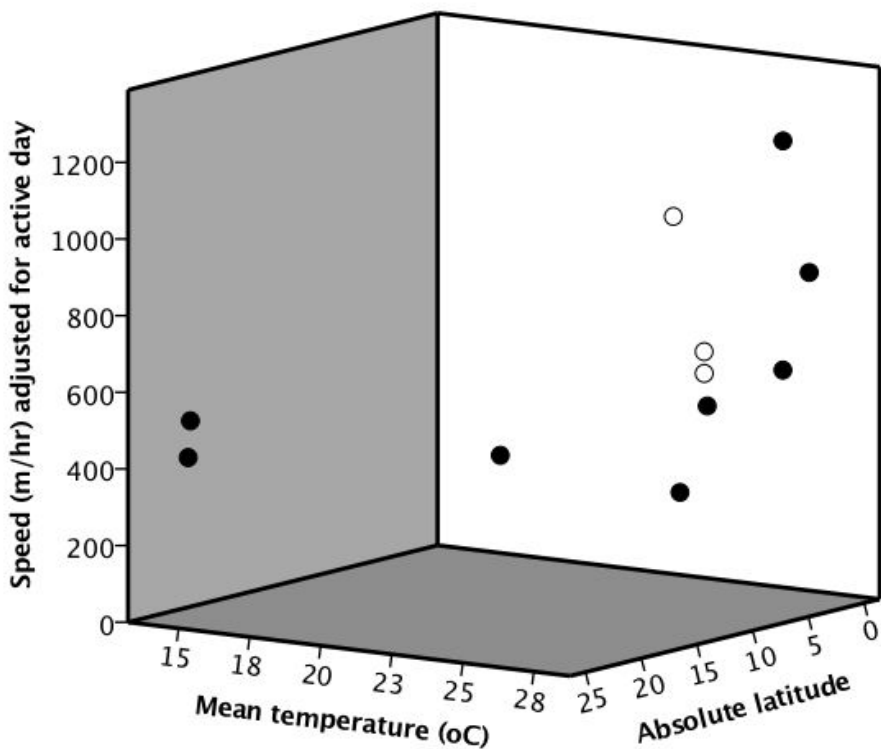
Mean speed for individual gibbon (filled symbols) and siamang (unfilled symbols) plotted against mean temperature and absolute latitude of the study site, assuming a 12-hour tropical day.

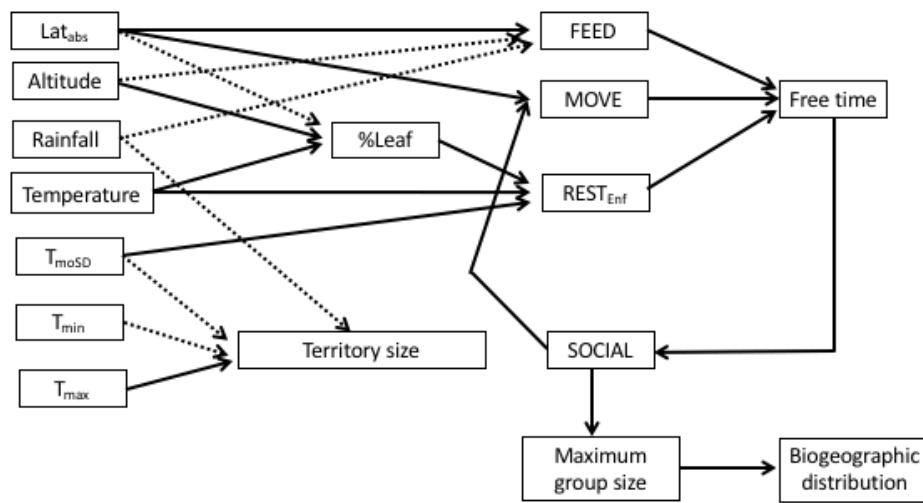


Many gibbon populations retire to nest several hours before sunset, with length of active daylength best predicted by mean rainfall (Table 2). To check whether a shorter active day makes any difference to these results, we recalculated speed as a function of actual length of active day using the equation given in Table 2 to predict length of active day from rainfall for each site. Aside from a slight uniform increase in speed, the transformation has no effect (Fig. S3b).

Figure S3(b)

Mean speed for individual gibbon (filled symbols) and siamang (unfilled symbols) plotted against mean temperature and absolute latitude of the study site, with speed calculated for active day length (estimated using the equation in rainfall from Table 2).





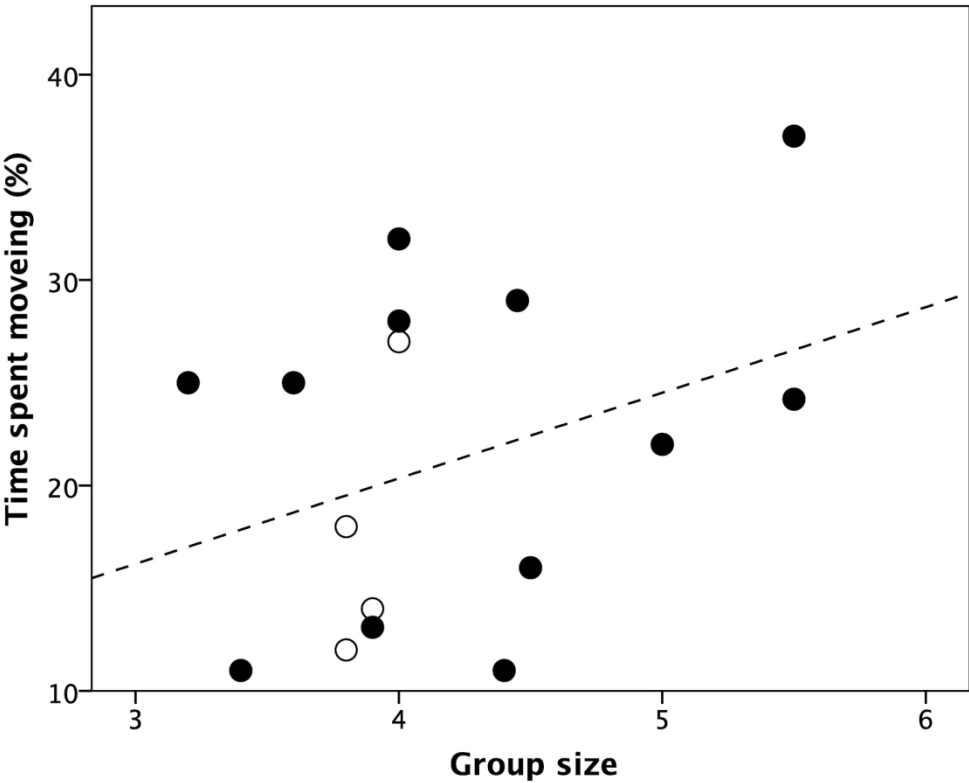


Fig. 2

165x131mm (300 x 300 DPI)

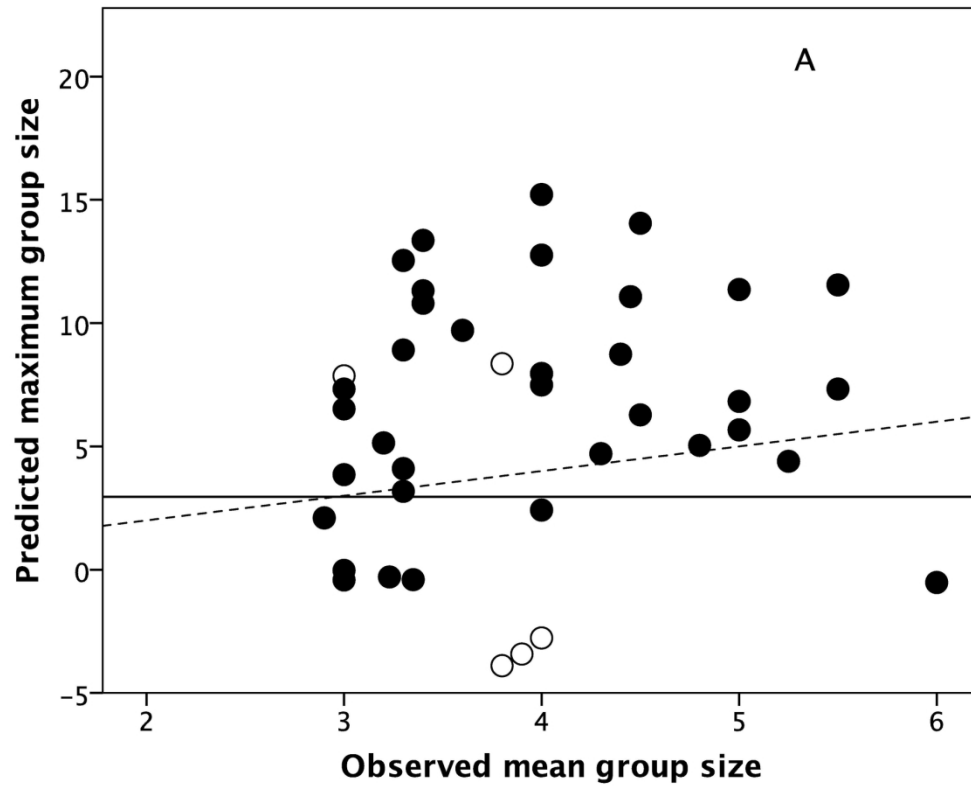


FIG. 3A

165x131mm (300 x 300 DPI)

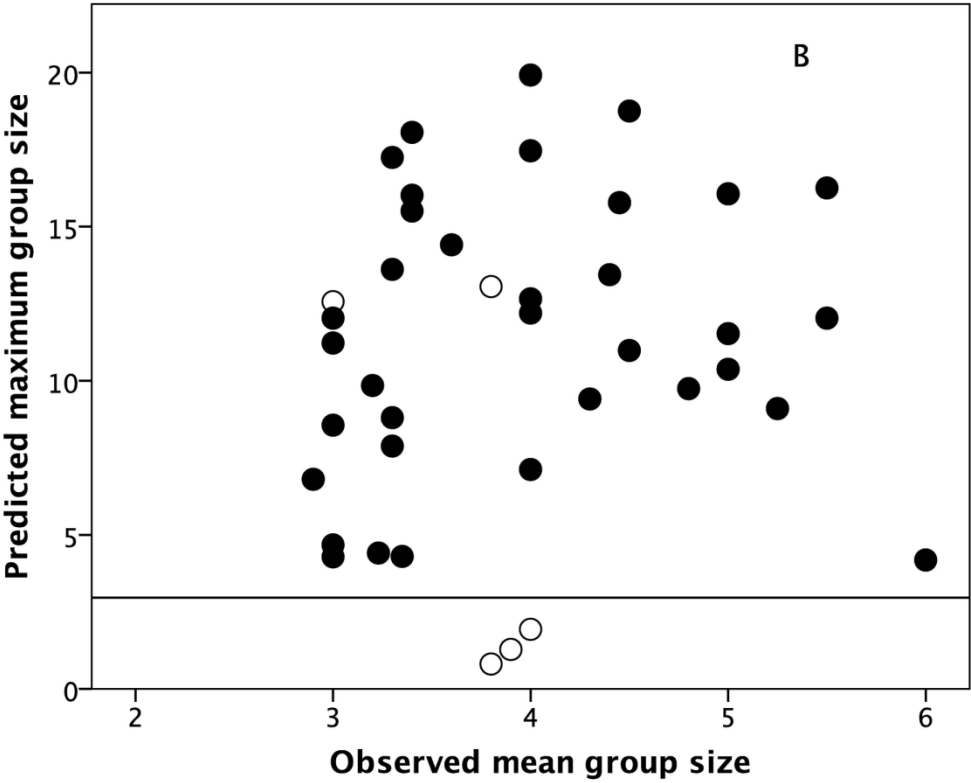


FIG. 3B

165x131mm (300 x 300 DPI)

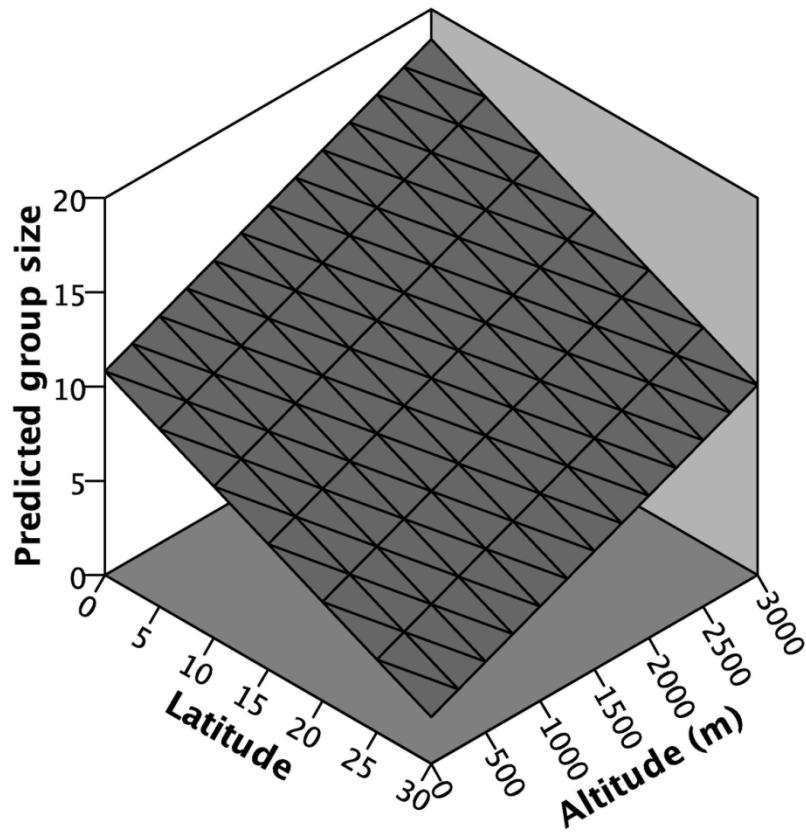


Fig. 4A

123x123mm (300 x 300 DPI)

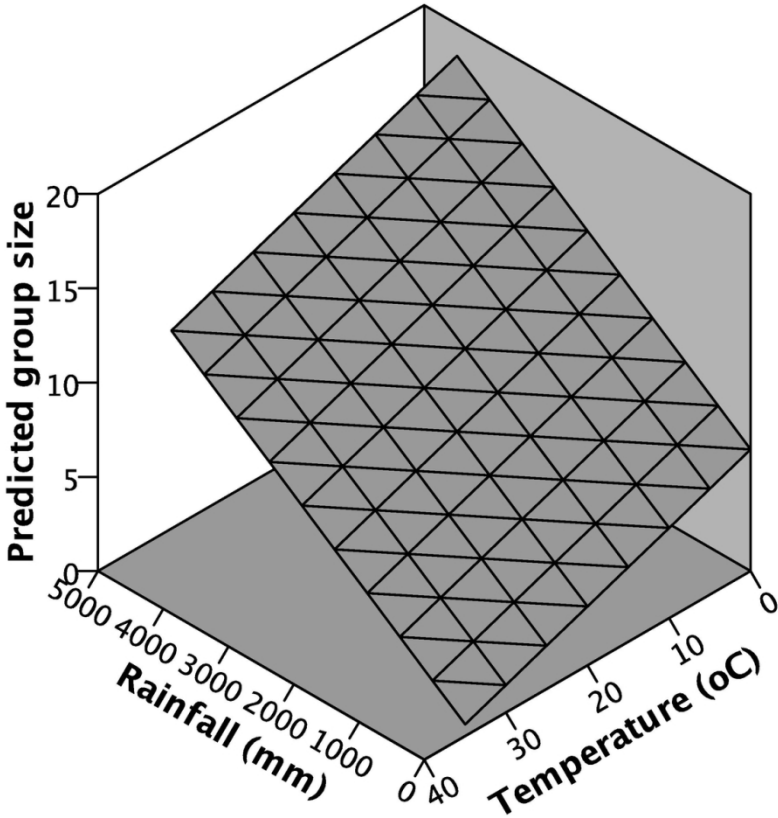


Fig. 4B

123x123mm (300 x 300 DPI)

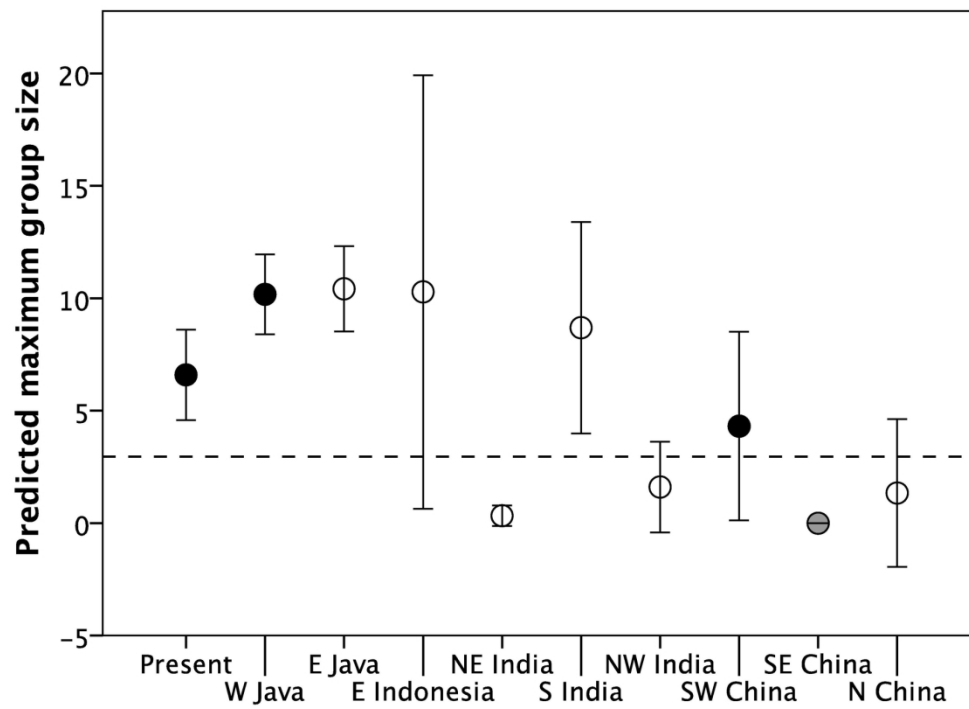


FIG. 5

165x131mm (300 x 300 DPI)

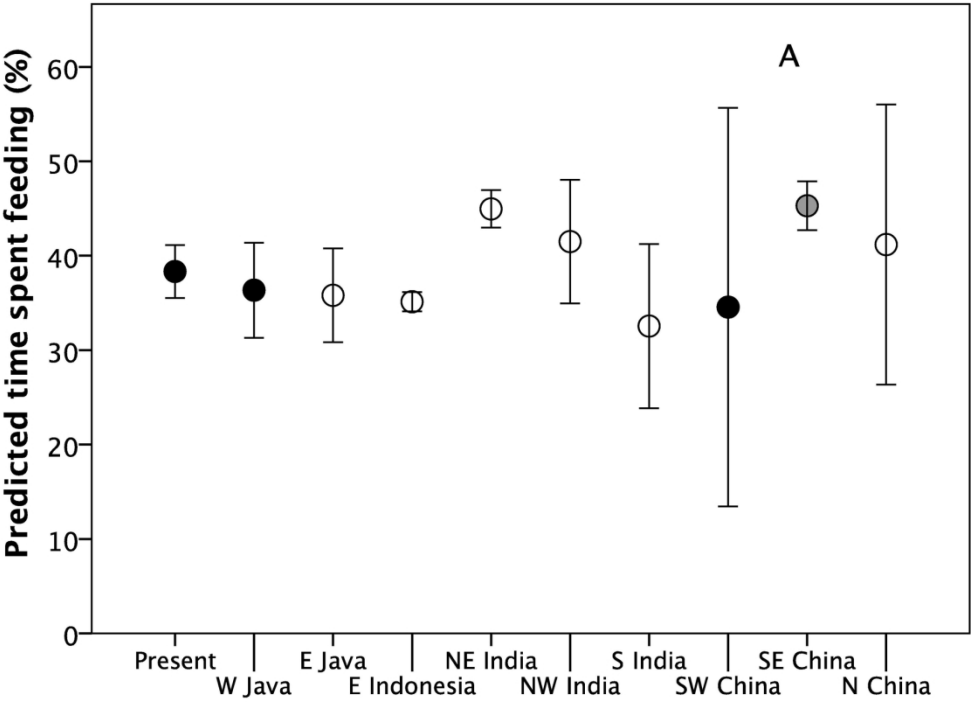


Fig. 6A

165x131mm (300 x 300 DPI)

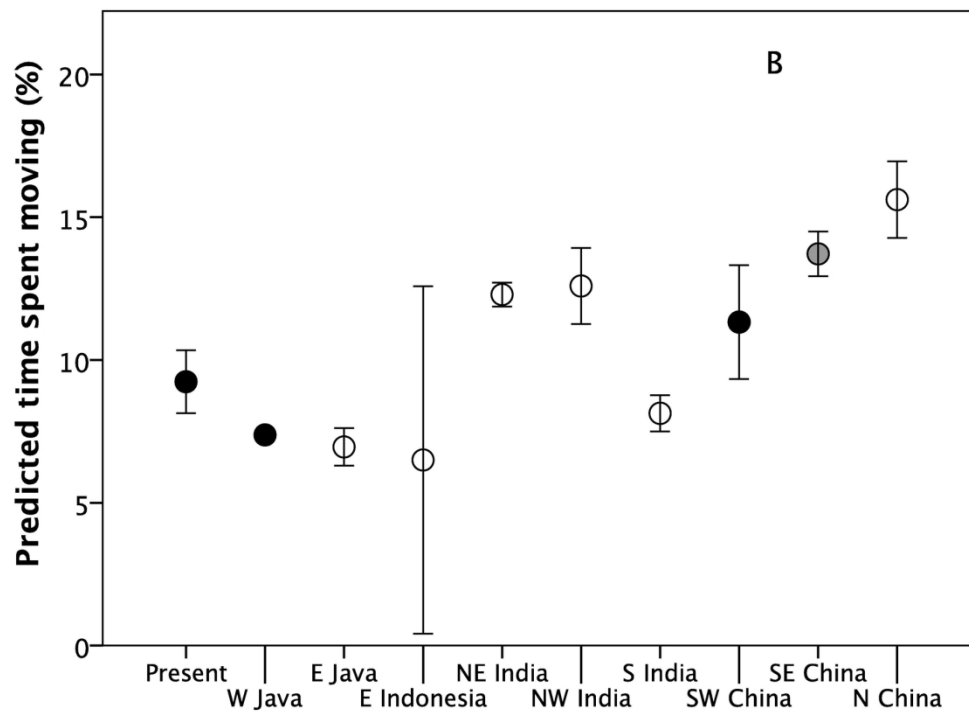


Fig. 6B

165x131mm (300 x 300 DPI)

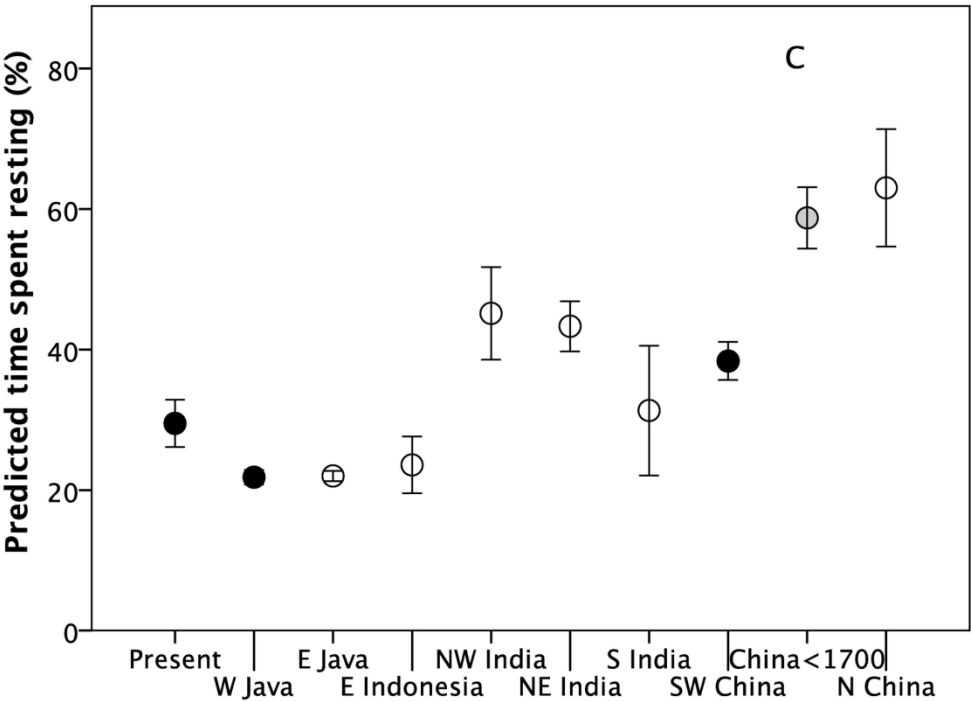


Fig. 6C

165x131mm (300 x 300 DPI)

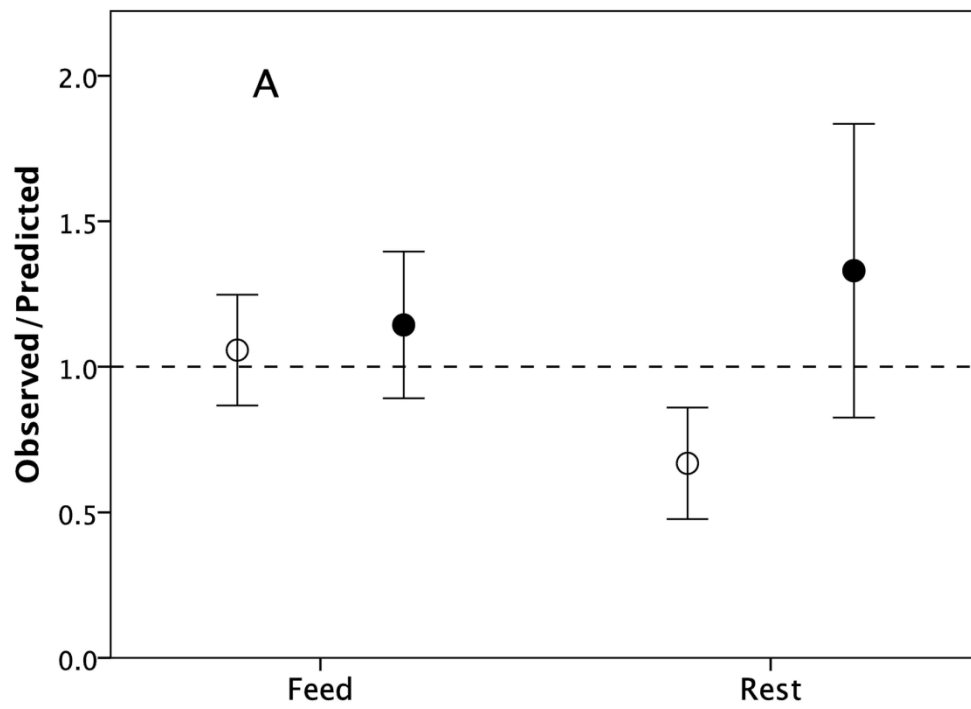


Fig. 7A

165x131mm (300 x 300 DPI)

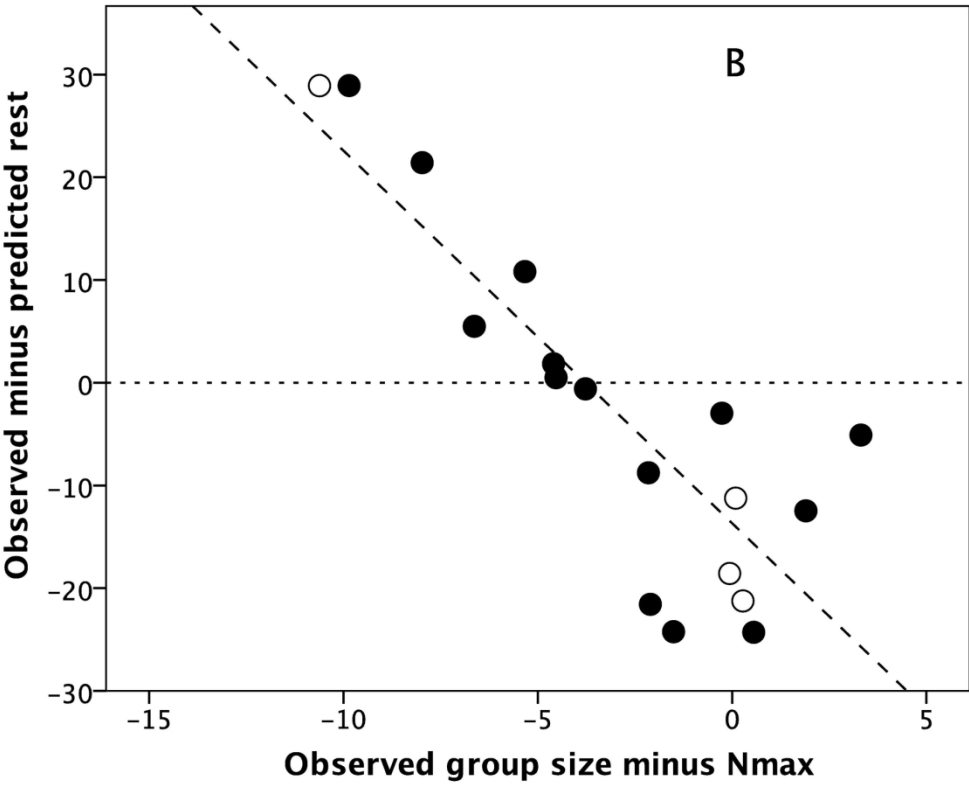


Fig. 7B

165x131mm (300 x 300 DPI)

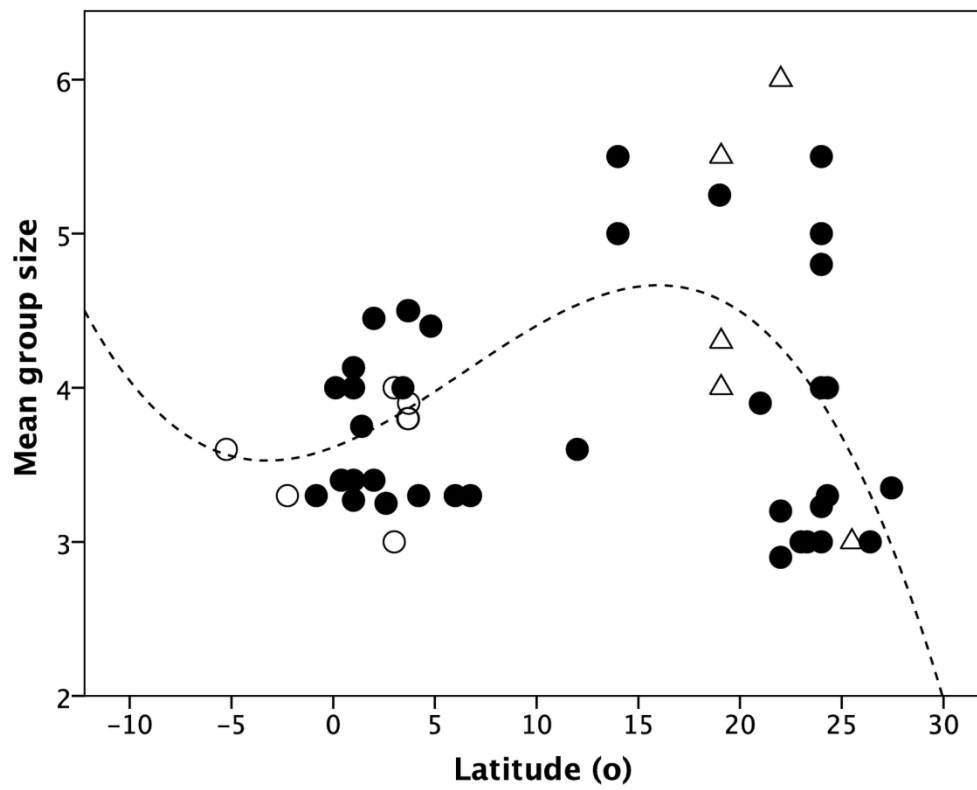


FIG. 8

165x131mm (300 x 300 DPI)